

GenCore version 5.1.5  
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OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 03:33:17 ; Search time 1431 Seconds  
(without alignments)  
1459.971 Million cell updates/sec

Title: US-09-689-430-1\_COPY\_150\_278

Perfect score: 129

Sequence: 1 cctcttaagtaaacagta.....gccatcagcgatcgatc 129

Scoring table: IDENTITY NUC  
Gapop 10.0, Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 32308132

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%  
Listing first 45 summaries

Database :

EST:\*  
1: em\_estdb:\*  
2: em\_esthum:\*  
3: em\_estin:\*  
4: em\_estnu:\*  
5: em\_estov:\*  
6: em\_estpl:\*  
7: em\_estro:\*  
8: em\_hlc:\*  
9: gb\_est1:\*  
10: gb\_est2:\*  
11: gb\_hlc:\*  
12: gb\_est3:\*  
13: gb\_est4:\*  
14: gb\_est5:\*  
15: em\_estfun:\*  
16: em\_estom:\*  
17: gb\_gss:\*  
18: em\_gss\_hum:\*  
19: em\_gss\_inv:\*  
20: em\_gss\_pln:\*  
21: em\_gss\_vit:\*  
22: em\_gss\_fun:\*  
23: em\_gss\_mam:\*  
24: em\_gss\_mus:\*  
25: em\_gss\_other:\*  
26: em\_gss\_pro:\*  
27: em\_gss\_rnd:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match Length	ID	Description
1	59.2	45.9	479	10 AV657611
2	46	35.7	451	10 AV685661
3	40.8	31.6	421	10 AV654002
4	32.8	25.4	927	14 B0642439
5	32.6	25.3	855	12 BF144380
6	32.2	25.0	466	9 AA616160

Result No.	Score	Query Match Length	ID	Description
7	31	24.0	915	13 B1561624
8	30.8	23.9	928	13 B1868197
9	30.6	23.7	213	10 AA445510
10	30.6	23.7	348	12 BF835894
11	30.6	23.7	377	12 BF835886
12	30.6	23.7	394	13 BM430717
13	30.6	23.7	458	13 BM429757
14	30.6	23.7	552	14 B0355187
15	30.6	23.7	567	14 AV596621
16	30.6	23.7	568	10 AV616893
17	30.6	23.7	750	10 BE296329
18	30.6	23.7	765	10 BE513735
19	30.6	23.7	915	14 B0648846
20	30.6	23.7	939	14 B0709796
21	30.6	23.7	951	14 B0709274
22	30.6	23.7	955	14 B0706617
23	30.6	23.7	1245	14 B066190
24	30.4	23.4	678	9 AL632211
25	30.2	23.3	678	9 AL632211
26	30	23.3	1063	10 AW953618
27	30	23.3	1063	12 BE871724
28	29.8	23.1	568	17 AF019127
29	29.6	22.9	519	17 A2902709
30	29.6	22.9	874	13 BM043559
31	29.6	22.9	888	13 BM011355
32	29.6	22.9	1060	13 BM470623
33	29.2	22.6	251	14 F05149
34	29.2	22.6	338	12 BF835885
35	29.2	22.6	462	12 BG956025
36	29.2	22.6	503	17 AQ853104
37	29.2	22.6	462	14 BE560380
38	29	22.5	462	14 BM967767
39	29	22.5	500	10 AW502350
40	29	22.5	743	12 BF862037
41	28.8	22.3	431	14 R89172
42	28.8	22.3	441	10 AV636155
43	28.8	22.3	547	10 AV619091
44	28.8	22.3	563	14 B0814899
45	28.8	22.3	663	14 B0824933

## ALIGNMENTS

### RESULT 1

LOCUS AV657611

DEFINITION AV657611 GLC Homo sapiens cDNA clone G1CFD601 3', mRNA sequence.

ACCESSION AV657611

VERSION AV657611.1 GI:9878625

KEYWORDS EST.

SOURCE human.

ORGANISM Homo sapiens

REFERENCE Xu,X., Huang,J., Xu,Z., Qian,B., Zhu,Z., Yan,Q., Cai,T., Zhang,X.,

AUTHORS Xieo,H., Qu,J., Liu,F., Huang,Q., Cheng,Z., Li,N., Du,J., Hu,W.,

TITLE Shen,K., Lu,G., Fu,G., Zhong,M., Xu,S., Gu,W., Huang,W., Zhao,X.,

INSIGHT INTO HEPATOCELLULAR CARCINOGENESIS AT TRANSCRIPTOME LEVEL

by comparing gene expression profiles of hepatocellular carcinoma

with those of corresponding noncancerous liver

Proc. Natl. Acad. Sci. U.S.A. 98 (26), 15089-15094 (2001)

21625106

CONTACT: Zeguang Han

Chinese National Human Genome Center at Shanghai

351 Guo Shoujing Road, Zhangjiang Hi-Tech Park, Pudong, Shanghai

201203, P. R. China

Tel: 86-21-50801919 (ex.45)

Fax: 86-21-50801922

Email: hanzg@chgc.sh.cn

This clone is available at CHGC in Shanghai.

```

FEATURES
  source
    Location/Qualifiers
      1..479
      /organism="Homo sapiens"
      /db_xref="taxon:9606"
      /clone="GICFDG01"
      /clone_1lb="GIC"
      /tissue_type="corresponding non cancerous liver tissue"
      /dev_stage="Adult"
      /lab_host="SOLR"
      /note="Vector: pBluescript sk(-); site_1: EcoRI; site_2:
      XhoI"
BASE COUNT      132 a      104 c      116 g      127 t
ORIGIN
Query Match      45.9%; Score 59.2; DB 10; Length 479;
Best Local Similarity 72.6%; Pred. No. 5.1e-09;
Matches 90; Conservative 0; Mismatches 33; Indels 1; Gaps 1;

QY 3 CTTTCTAGTAACAGTACATGACCTTTACCCGTTGCT-CGGCAACGGCCTGCTCTGT 61
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 263 CTGCTGAGGAACAATACATGACCTTTACCCGTTGATATGGCAACGGCAGTGTCTT 322
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

QY 62 GCCAAGTGTTCGTGACGACCCGACCTGGGCTTGCGCATATAGGCATACAGCGCA 121
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 323 CTCAGCTGTTCTCTATGCGTACACTAGTATGATGACAGATTGAACACGCTCTCATCAGCA 382
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

QY 122 TGGC 125
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 383 TGGC 386
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 2
LOCUS      AV685661      451 bp      mRNA      linear      EST 16-JAN-2002
DEFINITION AV685661 GKC Homo sapiens cDNA clone GKCCGH01 5', mRNA sequence.
ACCESSION  AV685661
VERSION     AV685661.1 GI:10287524
KEYWORDS   EST.
SOURCE      human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE  1 (bases 1 to 451)
            Xu,X., Huang,J., Xu,Z., Qian,B., Zhu,Z., Yan,Q., Cai,T., Zhang,X.,
            Xiao,H., Qu,J., Liu,F., Huang,Q., Cheng,Z., Li,N., Du,J., Hu,W.,
            Shen,K., Lu,G., Fu,G., Zhong,M., Xu,S., Gu,W., Huang,W., Zhao,X.,
            Hu,G., Gu,J., Chen,Z. and Han,Z.
            Insight into hepatocellular carcinogenesis at transcriptome level
            by comparing gene expression profiles of hepatocellular carcinoma
            with those of corresponding noncancerous liver
            Proc. Natl. Acad. Sci. U.S.A. 98 (26), 15089-15094 (2001)
            21625106
COMMENT     Contact: Zeguang Han
            Chinese National Human Genome Center at Shanghai
            351 Guo Shoujing Road, Zhangjiang Hi-Tech Park, Pudong, Shanghai
            201203, P. R. China
            Tel: 86-21-50801919(ex.45)
            Fax: 86-21-50801922
            Email: hanzge@chc.sh.cn
            This clone is available at CHGC in Shanghai.

FEATURES
  source
    Location/Qualifiers
      1..451
      /organism="Homo sapiens"
      /db_xref="taxon:9606"
      /clone="GKCCGH01"
      /clone_1lb="GKC"
      /tissue_type="hepatocellular carcinoma"
      /dev_stage="Adult"
      /lab_host="SOLR"
      /note="Vector: pBluescript sk(-); site_1: EcoRI; site_2:
      XhoI"
BASE COUNT      88 a      126 c      124 g      113 t
ORIGIN

```

```

Query Match      35.7%; Score 46; DB 10; Length 451;
Best Local Similarity 63.6%; Pred. No. 0.00012;
Matches 70; Conservative 0; Mismatches 40; Indels 0; Gaps 0;

QY 16 CAGTACATGACCTTTACCCCGTGTCTCGGCAACGGCCTGCTGTGCCAGTGTTCCT 75
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 1 CTTTAAGTCAACAATATACCGCCGTTGATCGCAATGCTAATCTGCAATATGATGATCT 60
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

QY 76 GACCAACCCCGACCTGGCTGGGCTGGCCATAGGCATACGCGATGCG 125
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 61 GACGCATACGCTACTGCTGCTGCTGGCCATGGCCATAGATCCGTGCG 110
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 3
LOCUS      AV654002      421 bp      mRNA      linear      EST 15-JAN-2002
DEFINITION AV654002 GKC Homo sapiens cDNA clone GICDQH06 3', mRNA sequence.
ACCESSION  AV654002
VERSION     AV654002.1 GI:9875016
KEYWORDS   EST.
SOURCE      human.
ORGANISM   Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE  1 (bases 1 to 421)
            Xu,X., Huang,J., Xu,Z., Qian,B., Zhu,Z., Yan,Q., Cai,T., Zhang,X.,
            Xiao,H., Qu,J., Liu,F., Huang,Q., Cheng,Z., Li,N., Du,J., Hu,W.,
            Shen,K., Lu,G., Fu,G., Zhong,M., Xu,S., Gu,W., Huang,W., Zhao,X.,
            Hu,G., Gu,J., Chen,Z. and Han,Z.
            Insight into hepatocellular carcinogenesis at transcriptome level
            by comparing gene expression profiles of hepatocellular carcinoma
            with those of corresponding noncancerous liver
            Proc. Natl. Acad. Sci. U.S.A. 98 (26), 15089-15094 (2001)
            21625106
COMMENT     Contact: Zeguang Han
            Chinese National Human Genome Center at Shanghai
            351 Guo Shoujing Road, Zhangjiang Hi-Tech Park, Pudong, Shanghai
            201203, P. R. China
            Tel: 86-21-50801919(ex.45)
            Fax: 86-21-50801922
            Email: hanzge@chc.sh.cn
            This clone is available at CHGC in Shanghai.

FEATURES
  source
    Location/Qualifiers
      1..421
      /organism="Homo sapiens"
      /db_xref="taxon:9606"
      /clone="GICDQH06"
      /clone_1lb="GIC"
      /tissue_type="corresponding non cancerous liver tissue"
      /dev_stage="Adult"
      /lab_host="SOLR"
      /note="Vector: pBluescript sk(-); site_1: EcoRI; site_2:
      XhoI"
BASE COUNT      112 a      99 c      82 g      128 t
ORIGIN
Query Match      31.6%; Score 40.8; DB 10; Length 421;
Best Local Similarity 77.5%; Pred. No. 0.0063;
Matches 62; Conservative 0; Mismatches 17; Indels 1; Gaps 1;

QY 3 CTTTCTAGTAACAGTACATGACCTT-TACCCGTTGCTGGCAACGGCCTGCTCTGT 61
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 342 CTTACTGCGTAACAATGCTAAATCTATATACCGGTTGACCGTACAGGATCATGTCTGT 401
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

QY 62 GCCAAGTGTTCGTGACGCA 81
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db 402 GCCAAGTGTTCGTGACGCA 421
    ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||

RESULT 4
LOCUS      B0642439      927 bp      mRNA      linear      EST 15-JUL-2002

```

DEFINITION AGENCOURT\_8286037 NIH\_MGC\_99 Homo sapiens cDNA clone IMAGE:6292602  
 5', mRNA sequence.  
 ACCESSION BO642439  
 VERSION BO642439.1 GI:21766611  
 EST.  
 SOURCE human.  
 ORGANISM Homo sapiens  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.  
 1 (bases 1 to 927)  
 NIH-MGC http://mgc.nci.nih.gov/.  
 National Institutes of Health, Mammalian Gene Collection (MGC)  
 UNPUBLISHED (1999)  
 COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: cga@rsf@mail.nih.gov  
 Tissue Procurement: Lou Staudt  
 cDNA Library Preparation: Rubin Laboratory  
 DNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Agencourt Bioscience Corporation  
 Clone distribution: MGC clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
 http://image.llnl.gov  
 Plate: LMC2494 row: e column: 19  
 High quality sequence stop: 539.  
 Location/Qualifiers  
 1..927  
 /organism="Homo sapiens"  
 /db\_xref="taxon:9606"  
 /clone\_image="6292602"  
 /clone\_id="NIH\_MGC\_99"  
 /tissue\_type="lymphoma, cell line"  
 /lab\_host="DH10B (phage-resistant)"  
 /note="Organ: lymph. Vector: pOTB7. Site 1: XhoI. Site 2:  
 EcoRI. cDNA made by oligo-dt priming. Directionally cloned  
 into EcoRI/XhoI sites using the following 5' adaptor:  
 GGCACGAG(g). Size-selected >500bp for average insert size  
 1.8kb. Library constructed by Ling Hong in the laboratory  
 of Gerald M. Rubin (University of California, Berkeley)  
 using ZAP-cDNA synthesis kit (Stratagene) and Superscript  
 II RT (Life Technologies). Note: this is a NIH\_MGC  
 library."  
 BASE COUNT 214 a 266 c 244 g 202 t 1 others  
 ORIGIN  
 Query Match 25.4%; Score 32.8; DB 14; Length 927;  
 Best Local Similarity 59.8%; Pred. No. 4.3;  
 Matches 55; Conservative 0; Mismatches 37; Indels 0; Gaps 0;  
 Oy 21 CATGACCTTACCCGCTGCTGCGCAACGGCGCTGCTGCGCAAGTGTTCGACGC 80  
 || ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||  
 Db 757 CAAACACCTTACACCTGAGAAAGGGGGGACCATCAAGCCCTTGCGCTTCAGCCCT 816  
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||  
 Oy 81 AACCCCACTGGCTGGGCTGGGCTGGCCATAGGCC 112  
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||  
 Db 817 AACCCCTACTGCTGGGCTGGGCTGGCCAGGCC 848  
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||  
 RESULT 5  
 BF144380/c 855 bp mRNA linear EST 24-OCT-2000  
 LOCUS 601787403F1 NCI\_CGAP\_Lu30 Mus musculus cDNA clone IMAGE:4015105 5',  
 mRNA sequence.  
 ACCESSION BF144380  
 VERSION BF144380.1 GI:10983420  
 EST.  
 SOURCE house mouse.  
 ORGANISM Mus musculus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 1 (bases 1 to 855)  
 NIH-MGC http://mgc.nci.nih.gov/.  
 National Institutes of Health, Mammalian Gene Collection (MGC)  
 UNPUBLISHED (1999)

COMMENT Contact: Robert Strausberg, Ph.D.  
 Email: cga@rsf@mail.nih.gov  
 Tissue Procurement: Gilbert Smith, Ph.D.  
 cDNA Library Preparation: Life Technologies, Inc.  
 DNA Library Arrayed by: The I.M.A.G.E. Consortium (LLNL)  
 DNA Sequencing by: Incyte Genomics, Inc.  
 Clone distribution: NCI-CGAP clone distribution information can be  
 found through the I.M.A.G.E. Consortium/LLNL at:  
 http://image.llnl.gov  
 Plate: LLM9260 row: f column: 02  
 High quality sequence stop: 693.  
 Location/Qualifiers  
 1..855  
 /organism="Mus musculus"  
 /strain="C57BL/6J"  
 /db\_xref="taxon:10090"  
 /clone\_image="4015105"  
 /clone\_id="NCI\_CGAP\_Lu30"  
 /tissue\_type="tumor, metastatic to mammary"  
 /lab\_host="DH10B"  
 /note="Organ: lung; Vector: pCMV-SPORT6; Site 1: NotI;  
 Site 2: SalI; Transgenic model MMTV-1, expression driven by  
 MMTV-LTR enhancer; Cloned unidirectionally. Primer: Oligo  
 dt. Library constructed by Life Technologies.  
 Investigator providing samples: Gilbert Smith, NIH"  
 BASE COUNT 188 a 211 c 279 g 177 t  
 ORIGIN  
 Query Match 25.3%; Score 32.6; DB 12; Length 855;  
 Best Local Similarity 69.8%; Pred. No. 4.8;  
 Matches 44; Conservative 0; Mismatches 19; Indels 0; Gaps 0;  
 Oy 1 CTCCTCTAGACAAACAGACATGACCTTACCCGCTGCGGACAGCGCTGCTG 60  
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||  
 Db 281 CTCCTCTAGACAAACAGACATGACCTTACCCGCTGCGGCTGCGGCTGCTG 222  
 ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||  
 Oy 61 TGC 63  
 |||||  
 Db 221 TGC 219  
 |||||  
 RESULT 6  
 AA616160/c 466 bp mRNA linear EST 07-OCT-1997  
 LOCUS v092905.1 B1 Barstead mouse irradiated colon MPEB7 Mus musculus cDNA  
 clone IMAGE:106616 5' similar to gb:M64716 405 RIBOSOMAL PROTEIN  
 S25 (HUMAN);, mRNA sequence.  
 ACCESSION AA616160  
 VERSION AA616160.1 GI:2503365  
 EST.  
 SOURCE house mouse.  
 ORGANISM Mus musculus  
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
 1 (bases 1 to 466)  
 Gelssel, S., Kucaba, T., Allen, M., Bowles, M., Dietrich, N., Dubnue, T.,  
 Schellenberg, K., Steptoe, M., Tan, F., Underwood, K., Moore, B.,  
 Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and  
 Waterston, R.  
 The WashU-HMI Mouse EST Project  
 UNPUBLISHED (1996)  
 COMMENT Contact: Marra M/Mouse EST Project  
 WashU-HMI Mouse EST Project  
 Washington University School of Medicine  
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
 Tel: 314 286 1800  
 Fax: 314 286 1810  
 Email: mouseest@wustl.edu  
 This clone is available royalty-free through LLNL; contact the  
 IMAGE Consortium (info@image.llnl.gov) for further information.  
 MGI:588976  
 Seq primer: -28m13 rev2 ET from Amersham

```

FEATURES                                High quality sequence stop: 107.
Source                                  Location/Qualifiers
1..466
/organism="Mus musculus"
/strain="FVB/N"
/db_xref="taxon:10090"
/clone="IMAGE:106616"
/clone_1lb="Barstead mouse irradiated colon MFLRB7"
/dev_stage="8 weeks"
/lab_host="DH10B"
(note="vector: pRT3D-Pac (Pharmacia) with a modified
polylinker; Site_1: EcoRI; Site_2: NotI; Tissue obtained
from 8 week old mouse. Colon was harvested 72 hours after
irradiation with 1400 Gys. 1st strand cDNA was primed
with a Not I - oligo(dT) primer
[5'GTTCAGATCTGAAGTGAGGACGGCGCCCTTTTCTTTTTTTTTTTTTTTT
T 3']; double-stranded cDNA was ligated to Eco RI
adaptors [AATTGCATCCTTG], digested with Not I and cloned
into the Not I and Eco RI sites of the modified pRT73
vector. Library constructed by Bob Barstead."
BASE COUNT      143 a          99 c       129 g        95 t
ORIGIN

Query Match                25.0%; Score 32.2; DB 9; Length 466;
Best Local Similarity     54.7%; Pred No. 4.8;
Matches    64; Conservative 0; Mismatches   53; Indels    0; Gaps    0;

Oy      2 TCTTTAGTAAGAAGTAGATGAACTTTACCCTGCTGCGCAAGCGCCTGCTGT 61
         ||| |||| | | | | | | | | | | | | | | | | | | | | |
Db      462 TTATCCAAAATTTCCACACTGTAAACCTTTTCACAAGAAATTTTCGACAGCATTTGGAGCGT 403

Oy      62 GCCAAGTGTTCCTGCACGCAACCCCACATCGCTGGGGCTTGCCCATAGGCCATCGAC 118
         |||| | ||| | | | | | | | | | | | | | | | | | | | | |
Db      402 CCCAACCCCTTGTGTCGCGGTGTAAATVAACTGGGGGCTGCTGCCCTTGGAACACAGC 346

RESULT 7
Bi561624/c
LOCUS      Bi561624              915 bp      mRNA      linear      EST 05-SEP-2001
DEFINITION 603256037/F1 NIH_MGC_97 Homo sapiens cDNA clone IMAGE:5298512 5',
            mRNA sequence.
VERSION     Bi561624
ACCESSION  Bi561624.1 GI:15448938
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE   1 (bases 1 to 915)
AUTHORS    NIH-MGC http://mgc.ncl.nih.gov/.
TITLE      National Institutes of Health, Mammalian Gene Collection (MGC)
JOURNAL    Unpublished (1999)
COMMENT     Contact: Robert Strausberg, Ph.D.
            Email: cgapbs@mail.nih.gov
            Tissue Procurement: Miklos Rakovits, M.D., Ph.D.
            CDNA Library Preparation: Michael J. Brownstein (NHGRI), Shitaki
            Toshiyuki and Piero Carninci (RIKEN)
            DNA sequencing by: Incyte Genomics, Inc.
            Clone distribution: MGC clone distribution information can be
            found through the I.M.A.G.E. Consortium/LNLM at:
            http://image.lnl.gov
            Plate: LHAM1755 row: 1 column: 09
            High quality sequence stop: 709.
            Location/Qualifiers
                1..915
                /organism="Homo sapiens"
                /db_xref="taxon:3606"
                /clone="IMAGE:5298512"
                /clone_1lb="NIH_MGC_97"
                /lab_host="DH10B"
                /note="Organ: testis; Vector: pBluescriptR (modified
                bluescript KS+); Site_1: BamHI; Site_2: SalI-XhoI (gtcgag

```

Query Match	Best Local Similarity	Score	DB	Length
Matches 67; Conservative	52.8%; Pred. No. 17;	24.0%;	DB 13;	915;
<p>); 01190-nt primed using primer 5'-TTTTTTTTTTTTTTTTTTVN-3' and size-selected for average insert size 2.2 kb and normalized to R0T 5. This is a primary library enriched for full-length clones and constructed using the Cap-trapper method (Carninci, in preparation). Library constructed by M. Brownstein (NIMH/NHGRI, National Institutes of Health). Note: this is a NIH_MGC Library."</p>				
BASE COUNT	248 a	192 c	257 g	218 t
ORIGIN				
Query Match	Best Local Similarity	Score <td>DB</td> <td>Length</td>	DB	Length
Matches 56; Conservative	57.1%; Pred. No. 20;	23.9%;	DB 13;	928;
<p>); 01190-nt primed using primer 5'-TTTTTTTTTTTTTTTTTTVN-3' and size-selected for average insert size 2.2 kb and normalized to R0T 5. This is a primary library enriched for full-length clones and constructed using the Cap-trapper method (Carninci, in preparation). Library constructed by M. Brownstein (NIMH/NHGRI, National Institutes of Health). Note: this is a NIH_MGC Library."</p>				
BASE COUNT	222 a	272 c	237 g	197 t
ORIGIN				

QY	21	CATGACACTTTTACCCCGTGTCTGGCAACGGCGCTGGTCTGCCAAGTGTTCGACGC	80
Db	738	CAAAACACCTTTACACCGCTGATGTGTGGGGAGATATCATCAACGCCCTGTGGCTTCACGCT	797
QY	81	AAACCCCACTGGCTGTGGGCTTGCCATAGGCCATCAGC	118
Db	798	AAACCGTACTGGCTGTGTCTGCCAGGAGCCAGCATC	835
RESULT 9			
LOCUS	AM445510/c	213 bp	mRNA linear
DEFINITION	812121 MARC 1BOV Bos taurus cDNA 5', mRNA sequence.		EST 25-APR-2001
ACCESSION	AM445510		
VERSION	AM445510.1	GI:6987272	
KEYWORDS	EST.		
SOURCE	cow.		
ORGANISM	Bos taurus		
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae; Bovidae; Bovinae; Bos.		
AUTHORS	1 (bases 1 to 213) Smith,T.P.L., Grosse,W.M., Fekling,B.A., Roberts,A.J., Stone,R.T., Casas,E., Wray,J.E., White,J., Cho,J., Fahrenkrug,S.C., Bennett,G.L., Heaton,M.P., Laegreid,W.M., Rohrer,G.A., Chitko-McKown,C.G., Pettes,G., Holt,I., Karamycheva,S., Liang,F., Quakenbush,J. and Keeler,J.W.		
TITLE	Sequence evaluation of four pooled-tissue normalized bovine cDNA libraries and construction of a gene index for cattle		
JOURNAL	Genome Res. 11 (4), 626-630 (2001)		
MEDLINE	21180013		
COMMENT	Contact: Smith TPL USDA, ARS, US Meat Animal Research Center PO Box 166, Clay Center, NE 68933-0166, USA Tel: 402 762 4366 Fax: 402 762 4390 Email: smith@email.marc.usda.gov Single pass sequencing. Bases called and trimmed with phred v0.980904.e. Vector identified by cross_match with the -mismscore 20 and -minmatch 12 options. PCR Primers FORWARD: AGGAACAGCTATGACCAT BACKWARD: GTTTCAGTCACGACG Plate: 43 row: A column: 20 Seq primer: ATTAGGTGACACTATAC. Location/Qualifiers 1. 213 /organism="Bos taurus" /db_xref="taxon:9913" /clone_lib="MARC 1BOV" /tissue_type="pooled" /lab_host="DH10B" /note="Vector: pCMV SPORT6; Site_1: NotI; Site_2: SalI; Library made from pooled tissue from lymph node, ovary, fat, hypothalamus, and pituitary."		
FEATURES			
source			
BASE COUNT	46 a 60 c	64 g 43 t	
ORIGIN			
Query Match	23.7%	Score 30.6;	DB 10; Length 213;
Best Local Similarity	60.0%	Pred. No. 11;	
Matches 51; Conservative	0;	Mismatches 34;	Indels 0; Gaps 0;
QY	11	GTAACACTTAATGATGAACTTTACCCCGTGTGGCGGACAGGCGTGTGTGGCCAAGTGT	70
Db	106	GTGACACACACAGACAGCACTTCGGCCACGATGCTCCAGAAAGGCGCAGAGGCGACCAAAAGCGG	47
QY	71	TTGTGACGCAACCCCACTGAGCTG	95
Db	46	TTGGCAGCGCAGCAAGCTGATGAGCTG	22
RESULT 10			

LOCUS	BPf835894	348 bp	mRNA	linear	EST 13-JAN-2001
DEFINITION	RC1-HT0975-161100-021-f05 HT0975 Homo sapiens CDNA, mRNA sequence.				
VERSION	BF835894				
KEYWORDS	BF835894.1	GI:12187336			
SOURCE	EST.				
ORGANISM	human.				
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.				
AUTHORS	Dias Neto, E., Garcia Correa, R., Verjovski-Almeida, S., Britones, M.R., Nagai, M.A., da Silva, W. Jr., Zago, M.A., Bordin, S., Costa, F.F., Goldstein, G.H., Carvalho, A.F., Matsukuma, A., Bala, G.S., Simpson, D.H., Brundman, A., deoliveira, P.S., Bucher, P., Jongeneel, C.V., O'Hare, M.J., Soares, F., Brentani, R.R., Reis, L.F., de Souza, S.J. and Simpson, A.J.				
TITLE	Shotgun sequencing of the human transcriptome with ORF expressed sequence tags				
JOURNAL	Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)				
MEDLINE	20202663				
COMMENT	Contact: Simpson A.J.G. Laboratory of Cancer Genetics Ludwig Institute for Cancer Research Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil Tel.: +55-11-2704922 Fax: +55-11-2707001 Email: asimpson@ludwig.org.br This sequence was derived from the RAPESP/LICR Human Cancer Genome Project. This entry can be seen in the following URL (http://www.ludwig.org.br/scripts/gethtml2.pl?rl=RC1c12-RC1-HT0975-161100-021-f05&t3=2000-11-16&t4=1) Seq primer: puc 18 forward High quality sequence stop: 348.				
FEATURES	Location/Qualifiers				
SOURCE	1..348 /organism="Homo sapiens" /db_xref="taxon:9606" /clone_1lb="HT0975" /dev_stage="Adult" /note="Organ: head_neck; Vector: puc18; Site:1: Sm1; Site:2: Sm1; A multi-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."				
BASE COUNT	65 a 102 c 125 g 56 t				
ORIGIN	Query Match 23.7%; Score 30.6; DB 12; Length 348; Best Local Similarity 53.8%; Pred. No. 14; Matches 63; Conservative 0; Mismatches 54; Indels 0; Gaps 0;				
QY	13 AAACGATCATGACCTTACCCGCTGCTCGGCAAGCGCTGCTGTGCCAAGTGT 72				
DB	162 AAAGATGAGTCGACACGCTGCTGAGGTGGCCGGGCCCTGTGAAAGCCTAGGGAT 221				
QY	73 GCTGACGCAACCCCACTGCGCTGGGGCTTGGCCATAGGCATACAGCGATGGGATC 129				
DB	222 GGAGAAAGTGAGCGACCGACGCGCTGGGGGCCCGCCGATAGCACAGCAGGCTCCGGGGGTC 278				
RESULT 11	BF835886 377 bp mRNA linear EST 13-JAN-2001				
LOCUS	BF835886				
DEFINITION	RC1-HT0975-161100-021-a09 HT0975 Homo sapiens CDNA, mRNA sequence.				
ACCESSION	BF835886				
VERSION	BF835886.1				
KEYWORDS	EST.				
SOURCE	human.				
ORGANISM	Homo sapiens				









JOURNAL	Arch. Virol. 143 (12), 2313-2326 (1998)
MEDLINE	99129050
REFERENCE	2 (pages 1 to 3221)
AUTHORS	Mishiro, S.
TITLE	Direct Submission
JOURNAL	Submitted (21-MAY-1998) Shunji Mishiro, Toshiba General Hospital, Dept. Medical Sciences; 6-3-22 Higashi Oh-1, Shinagawa, Tokyo 140 Japan (Tel:03-3764-8981, Fax:03-3764-8992)
FEATURES	
source	Location/Qualifiers 1..3221

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/organism="Hepatitis B virus"
/isolate="11D1HCC"
/db_xref="taxon:10407"
/note="genotype A-isolated from the circulation of a
japanese patient with hepatocellular carcinoma"
join(2307. .3221,1. .1623)
/gene="p"
join(2307. .3221,1. .1623)
CDS

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/protein_id="BA33871.1"
/db_xref="gi:3551317"
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LGNLWLSIPMTHAKVGNFTGLSVTPIENPMPQSPFNHILQEDILNRCCQFGPLTLP
VNEKRRLIPMTHAKVFNPHHTYLPLDKIKPYDOUVAVFORRHYLITLTKMGKILY
KRETTIRLSAFGSGPYSMEOELONGHRLVYKTSQRRDEFCGSPGJLILSRSSVGCITSS
LOKSRSLIPMTHAKVGLASQPSRGSTIARVYSPRRRTGYEFGSGGSHIDHSVNNSSS
CLHQSAVKRAVYSHLSTSRKSSSGCHAVEFICLPPNSAGSQSGQSSVSCMWLQDRNSK
PCESECLSHLNLREDMDPCDEHGHEHRIIRTPARVYGVFLVDKNEPHNTAEERLVVV
DEFQSPQRITRVSMKPAVNPVQSLTNLISLNLSTLSDVSAAYHILPLHPAARPHILL
IGSSGLITRVVARSNSNRILNNQGTQOMNLDDSCROLYSLMILLYKYGKRLHLYSEH
PIGGEFRITPMGVGLSPFLLOAFTSALCYVRRAAPHCLASVMDVDTLAKSVQJHVEH
ALYTPVTFNPLSLGSHLNPNTKRMGYSLNTMGITIGSWGILPDHDIYOKKHFPRKL
PVNRIIDNRKVCQRIYGLLGFAPFTQCYPALMPLRYATQAKAPFTSPYKALRSQ
YMNLYPVAKRQVGLQCVFADAPATPTGWMGLIGHQRNRPVAPLDIHTAEILADCFARS
SGGALITLSDNSVYLSRKYTSFPMPLGCAANNMILRGTSFYVPSALNPADDPSSGRGLG
LYRPLRLTRPEQPTGRTSLVAVSPSSVPHLPLVVRHVPASPLHVAHRPP"
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/gene="S"
/gene="S"
join(2854..3221,1..835)

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gene  
CDS

1374..1838  
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/codon\_start=1  
/product="x protein"  
/protein\_id="BA32869.1"  
/db\_xref="GI:3551315"

gene  
CDS

1374..1838  
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/codon\_start=1  
/product="x protein"  
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/db\_xref="GI:3551316"

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0y	3	CTTTCTAGTAACAGTACATGAACCTTTACCCCTTGCTCGCGAAGCGCCTGGTGTG	62
Db	1116	CTTTCTAGTAACAGTACATGAACCTTTACCCCTTGCTCGCGAAGCGCCTGGTGTG	1175
0y	63	CCAACTGTTTGGCTGACGCAACCCCGACGTCGTGGGGCTTGGCCATAGGCCATCAGCGCAT	122
Db	1176	CCAACTGTTTGGCTGACGCAACCCCGACGTCGTGGGGCTTGGCCATAGGCCATCAGCGCAT	1235
0y	123	GCG 125	
Db	1236	GCG 1238	

Search completed: May 21, 2003, 04:13:28  
Job time : 1028 secs

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VPADGHAHLISRLGIPCAFFSSGALCAFMETTVNHHQILPKVLRRTGLPA  
MSTIDLAVERDCEYKDEWELGEEIRLKFVILGCGRHLYCAAPAPCNFTSA"  
1814..2152  
/gene="C"  
1814..2152  
/gene="C"  
/note="pre-C, C"  
/codon\_start=1  
/product="truncated core antigen"  
/protein\_id="AA168823.1"  
/db\_xref="GI:1838998"  
/translation="MQLFHLCLISCTCPVQASKLCLGWLGMNDIDPYKEFGATVEL  
LSFLPSDFPFSVRLDITASALYREALSEPHCSPHHTALRQAILCWGELMTLATWVG  
NNLEDPHQIG"  
2136-2137  
/gene="C"  
/replace="a"  
BASE COUNT 746 a 864 c 707 g 903 t  
ORIGIN  
Query Match 95.3%; Score 123; DB 14; Length 3220;  
Best Local Similarity 100.0%; Pred. No. 4.4e-29;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 3 CTTTCTAAGTAACAGTACGACCTTTACCCCGTTGCTGGCAACGCGCTGCTGTG 62  
DB 1116 CTTTCTAAGTAACAGTACGACCTTTACCCCGTTGCTGGCAACGCGCTGCTGTG 1175  
QY 63 CCAAGTGTTCGTCAGCAACCCCACTGCTGGGCTTGGCCATAGGCGCATAGCGCAT 122  
DB 1176 CCAAGTGTTCGTCAGCAACCCCACTGCTGGGCTTGGCCATAGGCGCATAGCGCAT 1235  
QY 123 GCG 125  
DB 1236 GCG 1238  
RESULT 13  
AR085078 3221 bp DNA linear PAT 01-SEP-2000  
LOCUS AR085078  
DEFINITION Sequence 1 from patent US 5981274.  
ACCESSION AR085078  
VERSION AR085078.1 GI:10011849  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE 1 (bases 1 to 3221)  
AUTHORS Tyrrell,D.,Dorne,J., Chalsomchit,S. and Chang,L.-J.  
TITLE Recombinant hepatitis virus vectors  
JOURNAL Patent: US 5981274-A 1 09-NOV-1999;  
FEATURES  
Location/Qualifiers  
1..3221  
/organism="unknown"  
BASE COUNT 740 a 869 c 708 g 904 t  
ORIGIN  
Query Match 95.3%; Score 123; DB 6; Length 3221;  
Best Local Similarity 100.0%; Pred. No. 4.4e-29;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 3 CTTTCTAAGTAACAGTACGACCTTTACCCCGTTGCTGGCAACGCGCTGCTGTG 62  
DB 1118 CTTTCTAAGTAACAGTACGACCTTTACCCCGTTGCTGGCAACGCGCTGCTGTG 1177  
QY 63 CCAAGTGTTCGTCAGCAACCCCACTGCTGGGCTTGGCCATAGGCGCATAGCGCAT 122  
DB 1178 CCAAGTGTTCGTCAGCAACCCCACTGCTGGGCTTGGCCATAGGCGCATAGCGCAT 1237

QY 123 GCG 125  
DB 1238 GCG 1240  
RESULT 14  
E00010  
LOCUS E00010  
DEFINITION DNA of hepatitis B virus (HBV).  
ACCESSION E00010  
VERSION E00010.1 GI:2168319  
KEYWORDS JP 1981063995-A/1.  
SOURCE Hepatitis B virus.  
ORGANISM Hepatitis B virus  
REFERENCE 1 (bases 1 to 3221)  
AUTHORS Uchitama,T.R. and Hamada,M.G.  
TITLE NONTRANSIT VIRUS  
JOURNAL Patent: JP 1981063995-A 1 30-MAY-1981;  
COMMENT  
OS hepatitis B virus  
PN JP 1981063995-A/1  
PD 30-MAY-1981  
PF 24-MAY-1980 JP 1980069516  
PR 24-MAY-1979 US 79 41909, 26-DEC-1979 US 79 107267 PT  
URIMMU JIEE RATSUTAA, HAMADO MAIKERU GUTSUDOMAN PC  
C07H21/00,A61K39/29,C07C103/52,C07G7/00,C12N1/20,C12N15/00, PC  
C12P19/34//  
PC C12R1/19,C12R1/91;  
CC strandedness: double;  
CC topology: linear;  
CC hypothetical: No;  
CC anti-sense: No.  
FEATURES  
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/db\_xref="taxon:10407"  
BASE COUNT 740 a 872 c 705 g 904 t  
ORIGIN  
Query Match 95.3%; Score 123; DB 6; Length 3221;  
Best Local Similarity 100.0%; Pred. No. 4.4e-29;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 3 CTTTCTAAGTAACAGTACGACCTTTACCCCGTTGCTGGCAACGCGCTGCTGTG 62  
DB 1119 CTTTCTAAGTAACAGTACGACCTTTACCCCGTTGCTGGCAACGCGCTGCTGTG 1178  
QY 63 CCAAGTGTTCGTCAGCAACCCCACTGCTGGGCTTGGCCATAGGCGCATAGCGCAT 122  
DB 1179 CCAAGTGTTCGTCAGCAACCCCACTGCTGGGCTTGGCCATAGGCGCATAGCGCAT 1238  
QY 123 GCG 125  
DB 1239 GCG 1241  
RESULT 15  
AB014370  
LOCUS AB014370  
DEFINITION 3221 bp DNA circular VRL 06-FEB-1999  
ACCESSION AB014370  
VERSION AB014370.1 GI:3551314  
KEYWORDS  
SOURCE Hepatitis B virus (isolate:11D11HC) DNA.  
ORGANISM Hepatitis B virus  
REFERENCE 1 (sites)  
AUTHORS Takamashi,K., Akahane,Y., Hino,K., Ohta,Y. and Mishiro,S.  
TITLE Hepatitis B virus genomic sequence in the circulation of  
hepatocellular carcinoma patients: comparative analysis of 40  
full-length isolates

REMARK GenBank staff at the National Library of Medicine created this entry [NCBI gisbseq 138131] from the original journal article.

COMMENT This sequence comes from Fig. 5.

FEATURES Strong homology to human hepatitis B virus.

SOURCE Location/Qualifiers

1. 3218

/organism="Duck hepatitis B virus"

/specific\_host="duck"

/db\_xref="taxon:12639"

157..837

CDS

/gene="Pre-S/S"

/note="This sequence comes from Fig. 5"

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/product="Pre-S"

/protein\_id="A162458.1"

/db\_xref="GI:18158616"

/translation="MENITSGFLGPIVLAQGFLLTRITLITPOSIDSWTSTNIFLGGSPVCLGONSQSPSTNSPSCPPICPGYRMCLRRIFILFILLICILFLVLDYOGMLPVCPLIGTSTGCPCKTCTTPAGNSMPCSCCTKPSDNCICIPISSWAFAKYLWMAVSFRFMSLILVPEVQMFVGLSPVWLSAIIWMWYWGSPISYISPIPLPIFECILWYI"

157..834

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1376..1840

gene

/gene="X"

1376..1840

CDS

/gene="X"

/note="This sequence comes from Fig. 5"

/codon\_start=1

/product="X protein"

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complement(1622..2309)

gene

/gene="P"

/pseudo

/complement(1622..2309)

CDS

/gene="P"

/note="deletions at positions 2372 and 3028 resulting in stop codons and frame shift"

/pseudo

/codon\_start=1

1903..2475

gene

/gene="C"

/pseudo

1903..2475

CDS

/gene="C"

/note="stop codon within sequence"

/pseudo

/codon\_start=1

BASE COUNT 739 a 857 c 714 g 908 t

ORIGIN

Query Match 95.3%; Score 123; DB 14; Length 3218;

Best Local Similarity 100.0%; Pred. No. 4.4e-29;

Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 CTTTCTAGTAAACAGTACATGACCTTTACCCGTTGCTCGGCAAGCGCTGCTGTG 62

|||||

Db 1118 CTTTCTAGTAAACAGTACATGACCTTTACCCGTTGCTCGGCAAGCGCTGCTGTG 1177

QY 63 CCAAGTGTTCCTGACCAACCCCACTGGCTGGGCTTGCCATAGCCATCAGCGCAT 122

|||||

Db 1178 CCAAGTGTTCCTGACCAACCCCACTGGCTGGGCTTGCCATAGCCATCAGCGCAT 1237

QY 123 GCG 125

|||||

Db 1238 GCG 1240

RESULT 12

AF462041

LOCUS AF462041 3220 bp DNA Circular VRL 01-FEB-2002

DEFINITION Hepatitis B virus clone pAM6, complete genome.

ACCESSION AF462041

VERSION AF462041.1 GI:18389985

KEYWORDS

SOURCE

ORGANISM

REFERENCE

1 (bases 1 to 3220)

Authors Morlaty, A.M., Hoyer, B.H., Shih, J.W., Gerin, J.L. and Hamer, D.H.

Title Expression of the hepatitis B virus surface antigen gene in cell culture by using a simian virus 40 vector

Proc. Natl. Acad. Sci. U.S.A. 78 (4), 2606-2610 (1981)

JOURNAL MEDLINE 81233930

PUBMED 6264484

REFERENCE

2 (bases 1 to 3220)

Authors Jang, W.H., Yang, Y.I. and Kim, M.S.

Title Direct Submission

Submitted (20-DEC-2001) The Paik-Inje Memorial Institute for Biomedical Science, Inje University, 633-165 Gaeum-dong, Busanjin-gu, Busan 614-735, Korea

JOURNAL

FEATURES

SOURCE

Location/Qualifiers

1. 3220

/organism="Hepatitis B virus"

/db\_xref="ATCC:45020"

/db\_xref="taxon:10407"

/clone="pAM6: ATCC 45020"

/note="subtype: adw"

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/gene="P"

join(2306..3220,1..1623)

gene

/gene="P"

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/product="polymerase"

/protein\_id="A168823.1"

/db\_xref="GI:18389989"

/translation="MPLSYOHEFRKLLLDGTEAGPLLEELPRLADADLIRRYAEGLNGLANVSIPWTHKVGNETGLYSSTVPIFPEMOTPEPFRKIHLOEDILNKOOGVGLTVNEKRRKLIMPARFEPYTHKRYLPGLKGIKPYPDVVNHYVPTGRIYHIMKAGILYKRETTSAFSCGSPYSEDOELHNGRIYITRSQHDDESCSOSGLTSSSVPCIRS QKQSRGLGLOPHGPIASSOPGSRGSTRARVHPTIRCFEVEPSGSGHIDHSVNNSS CLHQSARAAVSHLSTSKQSSSGHAEVLCIPSSAGSOSGVSFCWMLQFRNSK PCSSEYCLSHLVNLRDEMGPCDEGEHHIIRTPPARVTCGVFLVDKNPHTAESRLVY DFESQFRGLTRVSWPKEFAVNPLOSLNLSNLSNLSLVSALFYIPIHPAMPHIL IGSSGLSRVYARLSNSRINNNQYGMONLHDCSRQIVYSALFYIKYGMKRLHYSH PLYLGRKIDMGVGLSPILLAOFTSAICSVPARAPHCACAFSTMDVYGAHSVORE SLTAVTNELSLGHLINPKTRKWTSLNFMGYVIGSKTLPDQHTVOKTRHCFKRL PVNRPIDMWVCORIVGLLGAPEFTQCGVPALMPLVACIOAKAFPSPTVYAFSKQ YNNLVPVARORPGLCQFADAPPTGGLAIGHQRMGTVEVAPLPIHTABLLAACFARS RSGAKLIGTDNSVYLSRKYTSFPMGLGTANMILRGTSFYVPSALNPADDDSRGLG LSRPLRLPPOPTGRTSLYAVGSPVSHLPVAVHVASPLHVAMRP"

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join(2853..3220,1..835)

CDS

/gene="S"

/note="pre-S1, pre-S2, S"

/codon\_start=1

/product="surface antigen"

/protein\_id="A168820.1"

/db\_xref="GI:1838986"

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1374..1838

gene

/gene="X"

1374..1838

CDS

/gene="X"

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OY 3 CTTTCTAGTAACAGTACATGACCTTTACCCCGTCTGCGCAACGCCCTGCTGTG 62
DB 2392 CTTTCTAGTAACAGTACATGACCTTTACCCCGTCTGCGCAACGCCCTGCTGTG 2451
OY 63 CCAAGTGTTCGTCAGCAGCAACCCCGACCTGGCGTGGCCATAGCCATACGCCAT 122
DB 2452 CCAAGTGTTCGTCAGCAGCAACCCCGACCTGGCGTGGCCATAGCCATACGCCAT 2511
OY 123 GCG 125
DB 2512 GCG 2514

RESULT 10
AF143306 3137 bp DNA circular VRL 19-OCT-1999
LOCUS Hepatitis B virus clone RM517, complete genome.
DEFINITION AF143306
ACCESSION AF143306.1 GI:5019974
KEYWORDS
SOURCE Hepatitis B virus.
ORGANISM Hepatitis B virus.
REFERENCE 1 (bases 1 to 3137)
AUTHORS Prekschat,P., Meisel,H., Will,H. and Gunther,S.
TITLE Hepatitis B virus genomes from long-term immunosuppressed virus
carriers are modified by specific mutations in several regions
JOURNAL J. Gen. Virol. 80 (Pt 10), 2685-2691 (1999)
MEDLINE 20037832
PUBMED 10573161

REFERENCE 2 (bases 1 to 3137)
AUTHORS Prekschat,P., Meisel,H., Iwanska,A., Will,H. and Gunther,S.
TITLE Direct Submission
JOURNAL Submitted (15-APR-1999) Department of Virology,
Bernhard-Nocht-Institute for Tropical Medicine,
Bernhard-Nocht-Strasse 74, Hamburg D-20359, Germany

FEATURES
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/organism="Hepatitis B virus"
/virus
/db_xref="taxon:10407"
/clone="RM517"
/note="amplified by PCR; contains a premature termination
codon in C gene, a deletion in pre-S/2 region removing
the pre-S2 start codon, a premature termination codon in S
gene, and a duplication plus insertion in X gene/core"
1. 171
/gene="pre-C/C"
1. 171
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/product="pre-C/C protein"
/protein_id="AAD37953.1"
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VERSION S65868.1 GI:436272
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SOURCE Duck hepatitis B virus.
ORGANISM Duck hepatitis B virus.
REFERENCE 1 (bases 1 to 3218)
AUTHORS Dai,W.L., Chen,Y., Li,L., Jiang,H.O. and Gu,J.R.
TITLE Nucleotide sequence of a cloned human HBV mutant (pDKHBV) in duck
hepatoma of Qidong County
JOURNAL Sci. China, Ser. B, Chem. Life Sci. Earth Sci. 36 (3), 329-338
(1993)
MEDLINE 94000357
PUBMED 8397804

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DB 2428 CTTTCTAGTAACAGTACATGACCTTTACCCCGTCTGCGCAACGCCCTGCTGTG 2487
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RESULT 11
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DEFINITION 3218 bp DNA linear VRL 16-JAN-2002
LOCUS S65868
DEFINITION Mutant, 4 genes, 3218 nt.
ACCESSION S65868
VERSION S65868.1 GI:436272
KEYWORDS
SOURCE Duck hepatitis B virus.
ORGANISM Duck hepatitis B virus.
REFERENCE 1 (bases 1 to 3218)
AUTHORS Dai,W.L., Chen,Y., Li,L., Jiang,H.O. and Gu,J.R.
TITLE Nucleotide sequence of a cloned human HBV mutant (pDKHBV) in duck
hepatoma of Qidong County
JOURNAL Sci. China, Ser. B, Chem. Life Sci. Earth Sci. 36 (3), 329-338
(1993)
MEDLINE 94000357
PUBMED 8397804

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GenCore version 5.1.5  
Copyright (c) 1993 - 2003 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 02:36:07 ; Search time 1023 Seconds  
(without alignments)  
3669.855 Million cell updates/sec

Title: US-09-689-430-1\_COPY\_150\_278

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Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 4109280

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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13: gb\_un.\*

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20: em\_om.\*

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22: em\_ov.\*

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38: em\_sy.\*

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40: em\_htgo\_mus.\*

41: em\_htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

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1	123	95.3	587	6	AR003584	AR003584 Sequence
2	123	95.3	587	6	AR062870	AR062870 Sequence
3	123	95.3	909	6	AR165345	AR165345 Sequence
4	123	95.3	1371	14	HBVNSAG2	M54898 Hepatitis B
5	123	95.3	2852	14	AF143307	AF143307 Hepatitis
6	123	95.3	3033	14	AF143299	AF143299 Hepatitis
7	123	95.3	3046	14	AF143301	AF143301 Hepatitis
8	123	95.3	3046	14	AF143308	AF143308 Hepatitis
9	123	95.3	3100	14	AF143305	AF143305 Hepatitis
10	123	95.3	3137	14	AF143306	AF143306 Hepatitis
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13	123	95.3	3221	6	AR085078	AR085078 Sequence
14	123	95.3	3221	6	E00010	E00010 DNA of hepa
15	123	95.3	3221	14	AB014370	AB014370 Hepatitis
16	123	95.3	3221	14	AB064314	AB064314 Hepatitis
17	123	95.3	3221	14	HBV012207	HBV012207 Hepatitis
18	123	95.3	3221	14	HBVADW2	X02763 Hepatitis B
19	123	95.3	3221	14	HBVXCPS	X70185 Hepatitis B
20	123	95.3	3221	14	HEB309370	AJ309370 Hepatitis
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26	123	95.3	4627	6	I03789	I03789 Sequence 4
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38	121.4	94.1	3161	14	AF143298	AF143298 Hepatitis
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# ALIGNMENTS

RESULT 1  
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LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

AR003584  
Sequence 1 from patent us 5744326.  
AR003584  
AR003584.1 GI:3964843

Unknown.  
Unclassified.

1 (bases 1 to 587)  
Ill.C.R. and Bidlingmaier,S.

Use of viral CIS-acting post-transcriptional regulatory sequences  
to increase expression of intronless genes containing  
near-consensus splice sites

587 bp  
DNA  
linear PAT 04-DEC-1998

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LOCUS AF143301
DEFINITION Hepatitis B virus clone BW1903, complete genome.
ACCESSION AF143301
VERSION AF143301.1 GI:5019947
KEYWORDS Hepatitis B virus.
SOURCE Hepatitis B virus
ORGANISM Hepatitis B virus
REFERENCE 1 (bases 1 to 3046)
AUTHORS Prekschat,P., Meisel,H., Will,H. and Gunther,S.
TITLE Hepatitis B virus genomes from long-term immunosuppressed virus
carriers are modified by specific mutations in several regions
JOURNAL J. gen. virol. 80 (Pt 10), 2685-2691 (1999)
MEDLINE 20037832
PUBMED 10573161
REFERENCE 2 (bases 1 to 3046)
AUTHORS Prekschat,P., Meisel,H., Iwanska,A., Will,H. and Gunther,S.
TITLE Direct Submission
JOURNAL Submitted (15-APR-1999) Department of Virology,
Bernhard-Nocht-Institute for Tropical Medicine,
Bernhard-Nocht-Strasse 74, Hamburg D-20359, Germany
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Db 2464 GCG 2466

RESULT 7
AF143301
LOCUS AF143301
DEFINITION Hepatitis B virus clone BW1903, complete genome.
ACCESSION AF143301
VERSION AF143301.1 GI:5019947
KEYWORDS Hepatitis B virus.
SOURCE Hepatitis B virus
ORGANISM Hepatitis B virus
REFERENCE 1 (bases 1 to 3046)
AUTHORS Prekschat,P., Meisel,H., Will,H. and Gunther,S.
TITLE Hepatitis B virus genomes from long-term immunosuppressed virus
carriers are modified by specific mutations in several regions
JOURNAL J. gen. virol. 80 (Pt 10), 2685-2691 (1999)
MEDLINE 20037832
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Bernhard-Nocht-Institute for Tropical Medicine,
Bernhard-Nocht-Strasse 74, Hamburg D-20359, Germany
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ACCESSION AF143301
VERSION AF143301.1 GI:5019947
KEYWORDS Hepatitis B virus.
SOURCE Hepatitis B virus
ORGANISM Hepatitis B virus
REFERENCE 1 (bases 1 to 3046)
AUTHORS Prekschat,P., Meisel,H., Will,H. and Gunther,S.
TITLE Hepatitis B virus genomes from long-term immunosuppressed virus
carriers are modified by specific mutations in several regions
JOURNAL J. gen. virol. 80 (Pt 10), 2685-2691 (1999)
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Bernhard-Nocht-Strasse 74, Hamburg D-20359, Germany
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RESULT 7
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LOCUS AF143301
DEFINITION Hepatitis B virus clone BW1903, complete genome.
ACCESSION AF143301
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KEYWORDS Hepatitis B virus.
SOURCE Hepatitis B virus
ORGANISM Hepatitis B virus
REFERENCE 1 (bases 1 to 3046)
AUTHORS Prekschat,P., Meisel,H., Will,H. and Gunther,S.
TITLE Hepatitis B virus genomes from long-term immunosuppressed virus
carriers are modified by specific mutations in several regions
JOURNAL J. gen. virol. 80 (Pt 10), 2685-2691 (1999)
MEDLINE 20037832
PUBMED 10573161
REFERENCE 2 (bases 1 to 3046)
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TITLE Direct Submission
JOURNAL Submitted (15-APR-1999) Department of Virology,
Bernhard-Nocht-Institute for Tropical Medicine,
Bernhard-Nocht-Strasse 74, Hamburg D-20359, Germany
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RESULT 7
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LOCUS AF143301
DEFINITION Hepatitis B virus clone BW1903, complete genome.
ACCESSION AF143301
VERSION AF143301.1 GI:5019947
KEYWORDS Hepatitis B virus.
SOURCE Hepatitis B virus
ORGANISM Hepatitis B virus
REFERENCE 1 (bases 1 to 3046)
AUTHORS Prekschat,P., Meisel,H., Will,H. and Gunther,S.
TITLE Hepatitis B virus genomes from long-term immunosuppressed virus
carriers are modified by specific mutations in several regions
JOURNAL J. gen. virol. 80 (Pt 10), 2685-2691 (1999)
MEDLINE 20037832
PUBMED 10573161
REFERENCE 2 (bases 1 to 3046)
AUTHORS Prekschat,P., Meisel,H., Iwanska,A., Will,H. and Gunther,S.
TITLE Direct Submission
JOURNAL Submitted (15-APR-1999) Department of Virology,
Bernhard-Nocht-Institute for Tropical Medicine,
Bernhard-Nocht-Strasse 74, Hamburg D-20359, Germany
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Best Local Similarity 100.0%; Pred. No. 4.4e-29;
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QY 123: GCG 125
Db 2449 GCG 2451
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LOCUS AF143308 3046 bp DNA circular VRL 19-OCT-1999
DEFINITION Hepatitis B virus clone WB1254, complete genome.
ACCESSION AF143308

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AF143308.1 GI:5019984  
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Hepatitis B virus  
Viruses; Retroviral viruses; Hepadnaviridae; Orthohepadnavirus.  
1 (bases 1 to 3046)  
Prekschat,P., Meisel,H., Will,H. and Gunther,S.  
Hepatitis B virus genomes from long-term immunosuppressed virus  
carriers are modified by specific mutations in several regions  
J. Gen. Virol. 80 (Pt 10), 2685-2691 (1999)  
20037832  
MEDLINE  
PUBMED 10573161  
REFERENCE 2 (bases 1 to 3046)  
Prekschat,P., Meisel,H., Iwanska,A., Will,H. and Gunther,S.  
Direct Submission  
TITLE  
Submitted (15-APR-1999) Department of Virology,  
Bernhard-Nocht-Institute for Tropical Medicine,  
Bernhard-Nocht-Strasse 74, Hamburg D-20359, Germany  
JOURNAL  
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Best Local Similarity 100.0%; Pred. No. 4.4e-29;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
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QY 63 CCAAGTGTGTTGCTGACCAACCCCACTGGCGTGGCGCATAGGCCATCAGCGCAT 122  
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QY 123 GCG 125  
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Db 2449 GCG 2451

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DEFINITION Hepatitis B virus clone RM501, complete genome.  
ACCESSION AF143305  
VERSION AF143305.1 GI:5019968  
KEYWORDS  
SOURCE Hepatitis B virus.  
ORGANISM Hepatitis B virus  
VIRUSES; Retroid viruses; Hepadnaviridae; Orthohepadnavirus.  
REFERENCE 1 (bases 1 to 3100)  
AUTHORS Preikschat,P., Meisel,H., Will,H. and Gunther,S.  
TITLE Hepatitis B virus genomes from long-term immunosuppressed virus  
carriers are modified by specific mutations in several regions  
J. Gen. Virol. 80 (Pt 10), 2685-2691 (1999)  
JOURNAL 20037832  
MEDLINE 10573161  
PUBMED  
REFERENCE 2 (bases 1 to 3100)  
AUTHORS Preikschat,P., Meisel,H., Iwanska,A., Will,H. and Gunther,S.  
TITLE Direct Submission  
Submitted (15-APR-1999) Department of Virology,  
Bernhard-Nocht-Institute for Tropical Medicine,  
Bernhard-Nocht-Strasse 74, Hamburg D-20359, Germany  
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Best Local Similarity 100.0%; Pred. No. 4.4e-29;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

GenCore version 5.1.5  
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OM nucleic - nucleic search, using sw model

Run on: May 21, 2003, 03:48:37 ; Search time 105 Seconds  
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1622.282 Million cell updates/sec

Title: US-09-689-430-1\_COPY\_150\_278

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Scoring table: IDENTITY NUC  
Gapop 10.0 , Gapext 1.0

Searched: 828747 seqs, 660231138 residues

Total number of hits satisfying chosen parameters: 1657494

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a  
score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

## SUMMARIES

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2	129	100.0	7944	12 US-10-095-718-1	Sequence 1, Appl1
3	119.8	92.9	3221	9 US-09-848-616-133	Sequence 133, App
4	110.2	85.4	306	9 US-09-875-453-25	Sequence 25, Appl
5	110.2	85.4	3182	9 US-10-104-966-14	Sequence 14, Appl
6	110.2	85.4	3182	10 US-09-929-955-14	Sequence 14, Appl
7	110.2	85.4	5618	9 US-10-142-358-1	Sequence 1, Appl1
8	110.2	85.4	7991	10 US-09-837-297-5	Sequence 5, Appl1
9	110.2	85.4	8007	10 US-09-837-297-3	Sequence 3, Appl1
10	110.2	85.4	8717	10 US-09-837-297-4	Sequence 4, Appl1
11	105	81.4	3215	9 US-10-209-264-1	Sequence 1, Appl1
12	99	76.7	5130	9 US-09-897-511A-9	Sequence 9, Appl1
13	99	76.7	5130	10 US-09-897-006-9	Sequence 9, Appl1
14	52.4	40.6	67	9 US-09-466-035-55	Sequence 55, Appl
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16	51.2	39.7	592	9 US-10-202-457-1	Sequence 1, Appl1
17	51.2	39.7	5691	9 US-09-897-511A-11	Sequence 11, Appl
18	51.2	39.7	5691	10 US-09-897-006-11	Sequence 11, Appl
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41	27.2	21.1	356	10 US-09-867-701-419	Sequence 253, App
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43	27	20.9	2714	9 US-10-227-884-21	Sequence 12, Appl
44	27	20.9	2714	9 US-10-230-163-21	Sequence 21, Appl
45	27	20.9	2714	9 US-10-218-631-21	Sequence 21, Appl

## ALIGNMENTS

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Sequence 3, Application US/10095718
Patent No. US20020131956A1
GENERAL INFORMATION:
APPLICANT: Walsh, Christopher
APPLICANT: Chao, Hengjun
APPLICANT: Burshtein, Haim
APPLICANT: Lynch, Carmel
APPLICANT: Stepan, Tony
APPLICANT: Munson, Keith
TITLE OF INVENTION: Adeno-Associated Virus Vectors Encoding Factor VIII and
FILE REFERENCE: 35052/204375
CURRENT APPLICATION NUMBER: US/10/095,718
CURRENT FILING DATE: 2002-03-12
PRIOR APPLICATION NUMBER: 09/689,430
PRIOR FILING DATE: 2001-08-22
PRIOR APPLICATION NUMBER: 60/158,780
PRIOR FILING DATE: 1999-10-12
NUMBER OF SEQ ID NOS: 5
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 3
LENGTH: 7914
TYPE: DNA
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: rAAV vector with canine B-domain deleted factor
FEATURE:
NAME/KEY: CDS
LOCATION: (435)...(4730)
US-10-095-718-3
Query Match 100.0%; Score 129; DB 12; Length 7914;
Best Local Similarity 100.0%; Pred. No. 8.5e-38;
Matches 129; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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DB 270 ATGCGGATC 278

## RESULT 2

US-10-095-718-1

Sequence 1, Application US/10095718  
Patent No. US20020131956A1  
GENERAL INFORMATION:  
APPLICANT: Walsh, Christopher  
APPLICANT: Chao, Hengjun  
APPLICANT: Burshtein, Haim  
APPLICANT: Lynch, Carmel  
APPLICANT: Stepan, Tony  
APPLICANT: Munson, Keith  
TITLE OF INVENTION: Adeno-Associated Virus Vectors Encoding Factor VIII and  
FILE REFERENCE: 35052/204375  
CURRENT APPLICATION NUMBER: US/10/095,718  
CURRENT FILING DATE: 2002-03-12  
PRIOR APPLICATION NUMBER: 09/689,430  
PRIOR FILING DATE: 2001-08-22  
PRIOR APPLICATION NUMBER: 60/158,780  
PRIOR FILING DATE: 1999-10-12  
NUMBER OF SEQ ID NOS: 5  
SOFTWARE: FastSeq for Windows Version 4.0  
SEQ ID NO 1  
LENGTH: 7944  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Plasmid pDLz6 encoding Homo sapiens BDD FVIII  
NAME/KEY: CDS  
LOCATION: (420)...(4835)  
US-10-095-718-1

Query Match 100.0%; Score 129; DB 12; Length 7944;  
Best Local Similarity 100.0%; Pred. No. 8.5e-38;  
Matches 129; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CTTCTTAAGTAACAGTACATGACCTTTACCCGCTGCTGGCAAGCGCTGGTCTG 60  
DB 150 CTTCTTAAGTAACAGTACATGACCTTTACCCGCTGCTGGCAAGCGCTGGTCTG 209  
QY 61 TGCCAAGTGTTCGTGACGCAACCCCACTGCTGGGGCTTGGCCATAGGCCATCAGCGC 120  
DB 210 TGCCAAGTGTTCGTGACGCAACCCCACTGCTGGGGCTTGGCCATAGGCCATCAGCGC 269  
QY 121 ATGCGGATC 129  
DB 270 ATGCGGATC 278

## RESULT 3

US-09-848-616-133

Sequence 133, Application US/09848616  
Publication No. US20030054010A1  
GENERAL INFORMATION:  
APPLICANT: Sebbel, Peter  
APPLICANT: Dunant, Nicolas  
APPLICANT: Bachmann, Martin  
APPLICANT: Tissot, Alain  
APPLICANT: Lechner, Franziska  
TITLE OF INVENTION: Molecular Antigen Array  
FILE REFERENCE: 1700.0180002  
CURRENT APPLICATION NUMBER: US/09/848,616  
CURRENT FILING DATE: 2001-05-05

NUMBER OF SEQ ID NOS: 186  
SOFTWARE: PatentIn Ver. 2.1  
SEQ ID NO 133  
LENGTH: 3221  
TYPE: DNA  
ORGANISM: Hepatitis B virus  
FEATURE:  
NAME/KEY: CDS  
LOCATION: (1901)..(2458)  
US-09-848-616-133

Query Match 92.9%; Score 119.8; DB 9; Length 3221;  
Best Local Similarity 98.4%; Pred. No. 1.7e-34;  
Matches 121; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 CTTCTTAAGTAACAGTACATGACCTTTACCCGCTGCTGGCAAGCGCTGGTCTG 62  
DB 1116 CTTCTTAAGTAACAGTACATGACCTTTACCCGCTGCTGGCAAGCGCTGGTCTG 1175  
QY 63 CCAAGTGTTCGTGACGCAACCCCACTGCTGGGGCTTGGCCATAGGCCATCAGCGCAT 122  
DB 1176 CCAAGTGTTCGTGACGCAACCCCACTGCTGGGGCTTGGCCATAGGCCATCAGCGCAT 1235  
QY 123 GCG 125  
DB 1236 GAG 1238

## RESULT 4

US-09-875-453-25

Sequence 25, Application US/09875453  
Publication No. US20030027320A1  
GENERAL INFORMATION:  
APPLICANT: Kim, Jungsub P.  
APPLICANT: Starr, Douglas B.  
APPLICANT: Tam, Albert W.  
APPLICANT: Laurance, Megan E.  
APPLICANT: Michelotti, Emil F.  
APPLICANT: Latour, Derek R.  
APPLICANT: Thomas, Rita L.  
APPLICANT: Kongsachith, Ana  
APPLICANT: Shepard, Liana T.  
APPLICANT: Lim, Moon Young  
APPLICANT: Brulice, Thomas W.  
TITLE OF INVENTION: PROMOTERS FOR REGULATED GENE EXPRESSION  
FILE REFERENCE: 4600-0135.30  
CURRENT APPLICATION NUMBER: US/09/875,453  
CURRENT FILING DATE: 2001-06-06  
PRIOR APPLICATION NUMBER: US 60/209,549  
PRIOR FILING DATE: 2000-06-06  
NUMBER OF SEQ ID NOS: 78  
SOFTWARE: FastSeq for Windows Version 4.0  
SEQ ID NO 25  
LENGTH: 306  
TYPE: DNA  
ORGANISM: Hepatitis B virus  
US-09-875-453-25

Query Match 85.4%; Score 110.2; DB 9; Length 306;  
Best Local Similarity 93.5%; Pred. No. 3.3e-31;  
Matches 115; Conservative 0; Mismatches 8; Indels 0; Gaps 0;

QY 3 CTTCTTAAGTAACAGTACATGACCTTTACCCGCTGCTGGCAAGCGCTGGTCTG 62  
DB 38 CTTCTTAAGTAACAGTACATGACCTTTACCCGCTGCTGGCAAGCGCTGGTCTG 97  
QY 63 CCAAGTGTTCGTGACGCAACCCCACTGCTGGGGCTTGGCCATAGGCCATCAGCGCAT 122  
DB 98 CCAAGTGTTCGTGACGCAACCCCACTGCTGGGGCTTGGCCATAGGCCATCAGCGCAT 157  
QY 123 GCG 125  
DB 1236 GAG 1238

US-09-929-955-14

SEQUENCE DESCRIPTION: SEQ ID NO: 1:  
5'-10-142-358-1

Query Match	85.4%;	Score 110.2;	DB 9;	Length 5618;
-------------	--------	--------------	-------	--------------

Best Local Similarity 93.5%; Pred. No. 7.4e-31;  
Matches 115; Conservative 0; Mismatches 8; Indels 0; Gaps 0;  
QY 3 CTTTGAAGTAACAGTACATGAACTTACCCCGTTGCTGCGCAAGCGCGCTGTGTG 62  
Db 1900 CTTTGTGTGTAACATACCTGAACTTACCCCGTTGCGCGCAAGCGCGCTGTGTG 1959  
QY 63 CCAAGTGTGTGCTGACGCAACCCCGCACTGCTGGGCTTGGCCATAGCCATAGCGCAT 122  
Db 1960 CCAAGTGTGTGCTGACGCAACCCCGCACTGCTGGGCTTGGCCATAGCGCAT 2019  
QY 123 GCG 125  
Db 2020 GCG 2022

RESULT 8  
US-09-837-297-5  
; Sequence 5, Application US/09837297  
; Patent No. US20010049145A1  
; GENERAL INFORMATION:  
; APPLICANT: RYU, WANG SHICK  
; TITLE OF INVENTION: Hepatitis B virus vectors for gene therapy  
; FILE REFERENCE:  
; CURRENT APPLICATION NUMBER: US/09/837,297  
; CURRENT FILING DATE: 2001-04-19  
; PRIOR APPLICATION NUMBER: KR2000-21070  
; PRIOR FILING DATE: 2000-04-20  
; NUMBER OF SEQ ID NOS: 5  
; SOFTWARE: Kopatentlin 1.71  
; SEQ ID NO 5  
; LENGTH: 7991  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: R712: PCMV-HBV/GFP3.2 Full Sequence  
US-09-837-297-5

Query Match 85.4%; Score 110.2; DB 10; Length 7991;  
Best Local Similarity 93.5%; Pred. No. 8.1e-31;  
Matches 115; Conservative 0; Mismatches 8; Indels 0; Gaps 0;  
QY 3 CTTTGAAGTAACAGTACATGAACTTACCCCGTTGCTGCGCAAGCGCGCTGTGTG 62  
Db 2465 CTTTGTGTGTAACATACCTGAACTTACCCCGTTGCGCGCAAGCGCGCTGTGTG 2524  
QY 63 CCAAGTGTGTGCTGACGCAACCCCGCACTGCTGGGCTTGGCCATAGCCATAGCGCAT 122  
Db 2525 CCAAGTGTGTGCTGACGCAACCCCGCACTGCTGGGCTTGGCCATAGCGCAT 2584  
QY 123 GCG 125  
Db 2585 GCG 2587

RESULT 9  
US-09-837-297-3  
; Sequence 3, Application US/09837297  
; Patent No. US20010049145A1  
; GENERAL INFORMATION:  
; APPLICANT: RYU, WANG SHICK  
; TITLE OF INVENTION: Hepatitis B virus vectors for gene therapy  
; FILE REFERENCE:  
; CURRENT APPLICATION NUMBER: US/09/837,297  
; CURRENT FILING DATE: 2001-04-19  
; PRIOR APPLICATION NUMBER: KR2000-21070  
; PRIOR FILING DATE: 2000-04-20  
; NUMBER OF SEQ ID NOS: 5  
; SOFTWARE: Kopatentlin 1.71  
; SEQ ID NO 3  
; LENGTH: 8007  
; TYPE: DNA  
; ORGANISM: HBV

FEATURE:  
; NAME/KEY: gene  
; LOCATION: (1)..(8007)  
; OTHER INFORMATION: Prototype vector of HBV  
US-09-837-297-3

Query Match 85.4%; Score 110.2; DB 10; Length 8007;  
Best Local Similarity 93.5%; Pred. No. 8.1e-31;  
Matches 115; Conservative 0; Mismatches 8; Indels 0; Gaps 0;  
QY 3 CTTTGAAGTAACAGTACATGAACTTACCCCGTTGCTGCGCAAGCGCGCTGTGTG 62  
Db 2481 CTTTGTGTGTAACATACCTGAACTTACCCCGTTGCGCGCAAGCGCGCTGTGTG 2540  
QY 63 CCAAGTGTGTGCTGACGCAACCCCGCACTGCTGGGCTTGGCCATAGCCATAGCGCAT 122  
Db 2541 CCAAGTGTGTGCTGACGCAACCCCGCACTGCTGGGCTTGGCCATAGCGCAT 2600  
QY 123 GCG 125  
Db 2601 GCG 2603

RESULT 10  
US-09-837-297-4  
; Sequence 4, Application US/09837297  
; Patent No. US20010049145A1  
; GENERAL INFORMATION:  
; APPLICANT: RYU, WANG SHICK  
; TITLE OF INVENTION: Hepatitis B virus vectors for gene therapy  
; FILE REFERENCE:  
; CURRENT APPLICATION NUMBER: US/09/837,297  
; CURRENT FILING DATE: 2001-04-19  
; PRIOR APPLICATION NUMBER: KR2000-21070  
; PRIOR FILING DATE: 2000-04-20  
; NUMBER OF SEQ ID NOS: 5  
; SOFTWARE: Kopatentlin 1.71  
; SEQ ID NO 4  
; LENGTH: 8717  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: R711: PCMV-HBV/GFP Full Sequence  
US-09-837-297-4

Query Match 85.4%; Score 110.2; DB 10; Length 8717;  
Best Local Similarity 93.5%; Pred. No. 8.3e-31;  
Matches 115; Conservative 0; Mismatches 8; Indels 0; Gaps 0;  
QY 3 CTTTGAAGTAACAGTACATGAACTTACCCCGTTGCTGCGCAAGCGCGCTGTGTG 62  
Db 3191 CTTTGTGTGTAACATACCTGAACTTACCCCGTTGCGCGCAAGCGCGCTGTGTG 3250  
QY 63 CCAAGTGTGTGCTGACGCAACCCCGCACTGCTGGGCTTGGCCATAGCCATAGCGCAT 122  
Db 3251 CCAAGTGTGTGCTGACGCAACCCCGCACTGCTGGGCTTGGCCATAGCGCAT 3310  
QY 123 GCG 125  
Db 3311 GCG 3313

RESULT 11  
US-10-209-264-1  
; Sequence 1, Application US/10209264  
; Publication No. US20030003111A1  
; GENERAL INFORMATION:  
; APPLICANT: Oon, Chong Jin  
; Lhm, Gek Keow  
; Zhao, Yi  
; Chen, Wei Ning  
; TITLE OF INVENTION: A MUTANT HUMAN HEPATITIS B VIRAL STRAIN AND  
; USES THEREOF

NUMBER OF SEQUENCES: 11  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Ladas & Parry  
STREET: 26 West 61 Street  
CITY: New York  
STATE: New York  
COUNTRY: USA  
ZIP: 10023  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.30  
CURRENT APPLICATION NUMBER: US/10/209,264  
FILING DATE: 31-Jul-2002  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: PCT/SG98/00046  
FILING DATE: 19-JAN-1998  
ATTORNEY/AGENT INFORMATION:  
NAME: Mass, Clifford J.  
REGISTRATION NUMBER: 30,086  
REFERENCE/DOCKET NUMBER: U-013109-7  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (212) 708-1800  
INFORMATION FOR SEQ ID NO: 1:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 3215 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: circular  
SEQUENCE DESCRIPTION: SEQ ID NO: 1:  
US-10-209-264-1

Query Match 81.4%; Score 105; DB 9; Length 3215;  
Best Local Similarity 91.7%; Pred. No. 5.4e-29;  
Matches 111; Conservative 0; Mismatches 10; Indels 0; Gaps 0;

3 CTTTCTAGTAAAGTACATGACCTTTACCCGTTGCTGCGCAACGGCTGCTGTG 62  
DB 1116 CTTTCTGTGTAACATATCTGACCTTTACCCGTTGCGCAACGGCTGCTGTG 1175

63 CCAAGTGTGTTGCTGACGCAACCCCACTGCTGGGCTTGCCATAGCCATCAGCGCAT 122  
DB 1176 CCAAGTGTGTTGCTGACGCAACCCCACTGATGGGCTTGCCATAGCCATCAGCGCAT 1235

123 G 123  
DB 1236 G 1236

RESULT 12  
US-09-897-511A-9  
Sequence 9, Application US/09897511A  
Publication No. US20030092882A1  
GENERAL INFORMATION:  
APPLICANT: Bremel, Robert  
APPLICANT: Miller, Linda  
TITLE OF INVENTION: Host Cells Containing Multiple Integrating Vectors  
FILE REFERENCE: GALA-06416  
CURRENT APPLICATION NUMBER: US/09/897,511A  
CURRENT FILING DATE: 2001-06-29  
PRIOR APPLICATION NUMBER: 60/215,925  
PRIOR FILING DATE: 2000-07-03  
NUMBER OF SEQ ID NOS: 36  
SOFTWARE: Patentin version 3.0  
SEQ ID NO 9  
LENGTH: 5130  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:

OTHER INFORMATION: Synthetic  
US-09-897-511A-9

Query Match 76.7%; Score 99; DB 9; Length 5130;  
Best Local Similarity 87.8%; Pred. No. 1e-26;  
Matches 108; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

3 CTTTCTAGTAAAGTACATGACCTTTACCCGTTGCTGCGCAACGGCTGCTGTG 62  
DB 1995 CTTTCTGTGTAACATATCTGACCTTTACCCGTTGCGCAACGGCTGCTGTG 2054

63 CCAAGTGTGTTGCTGACGCAACCCCACTGCTGGGCTTGCCATAGCCATCAGCGCAT 122  
DB 2055 CCAAGTGTGTTGCTGACGCAACCCCACTGATGGGCTTGCCATAGCCATCAGCGCAT 2114

123 GCG 125  
DB 2115 GCG 2117

RESULT 13  
US-09-897-006-9  
Sequence 9, Application US/09897006  
Patent No. US20020106729A1  
GENERAL INFORMATION:  
APPLICANT: Black, Gregory  
TITLE OF INVENTION: Expression Vectors  
FILE REFERENCE: GALA-06415  
CURRENT APPLICATION NUMBER: US/09/897,006  
CURRENT FILING DATE: 2001-06-29  
PRIOR APPLICATION NUMBER: 60/215,851  
PRIOR FILING DATE: 2000-07-03  
NUMBER OF SEQ ID NOS: 36  
SOFTWARE: Patentin version 3.0  
SEQ ID NO 9  
LENGTH: 5130  
TYPE: DNA  
ORGANISM: Artificial Sequence  
FEATURE:  
OTHER INFORMATION: Synthetic  
US-09-897-006-9

Query Match 76.7%; Score 99; DB 10; Length 5130;  
Best Local Similarity 87.8%; Pred. No. 1e-26;  
Matches 108; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

3 CTTTCTAGTAAAGTACATGACCTTTACCCGTTGCTGCGCAACGGCTGCTGTG 62  
DB 1995 CTTTCTGTGTAACATATCTGACCTTTACCCGTTGCGCAACGGCTGCTGTG 2054

63 CCAAGTGTGTTGCTGACGCAACCCCACTGCTGGGCTTGCCATAGCCATCAGCGCAT 122  
DB 2055 CCAAGTGTGTTGCTGACGCAACCCCACTGATGGGCTTGCCATAGCCATCAGCGCAT 2114

123 GCG 125  
DB 2115 GCG 2117

RESULT 14  
US-09-466-035-55  
Sequence 55, Application US/09466035  
Patent No. US20020165172A1  
GENERAL INFORMATION:  
APPLICANT: SALBERG, MATTI  
MILICH, DAVID R.  
LEE, WILLIAM T. L.  
TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR TREATING INTRACELLULAR DISEASES  
NUMBER OF SEQUENCES: 86  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Robins & Pasternak LLP  
STREET: 545 Middlefield Road, Suite 180

CITY: Menlo Park  
STATE: California  
COUNTRY: U.S.  
ZIP: 94025  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/466,035  
FILING DATE: 17-Dec-1999  
CLASSIFICATION: <Unknown>  
ATTORNEY/AGENT INFORMATION:  
NAME: Pasternak, Darna S.  
REGISTRATION NUMBER: 41,411  
REFERENCE/DOCKET NUMBER: 2300-1231.01  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 650-325-7812  
TELEFAX: 650-325-7823  
TELEX: <Unknown>  
INFORMATION FOR SEQ ID NO: 55:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 67 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
SEQUENCE DESCRIPTION: SEQ ID NO: 55:  
US-09-466-035-55

Query Match 40.6%; Score 52.4; DB 9; Length 67;  
Best Local Similarity 98.1%; Pred. No. 6.2e-10;  
Matches 53; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 72 TCGTGACGCAACCCCACTGCTGGGGCTTGSCCATAGGCAATCAGCGCATGCG 125  
|||  
DB 14 TCGTGACGCAACCCCACTGCTGGGGCTTGSCCATAGGCAATCAGCGCATGCG 67

RESULT 15  
US-09-912-679-55  
Sequence 55, Application US/09912679  
Patent No. US20020141974A1  
GENERAL INFORMATION:  
APPLICANT: Jolly, Douglas J.  
Chang, Stephen M.W.  
Lee, William T.L.  
Townsend, Kay  
O'Dea, Joanne  
TITLE OF INVENTION: HEPATITIS THERAPEUTICS  
NUMBER OF SEQUENCES: 84  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Seed and Berry  
STREET: 6300 Columbia Center, 701 Fifth Avenue  
CITY: Seattle  
STATE: Washington  
COUNTRY: U.S.  
ZIP: 98104  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/912,679  
FILING DATE: 07-Jun-1995  
CLASSIFICATION: <Unknown>  
ATTORNEY/AGENT INFORMATION:  
NAME: Mcmasters, David D.  
REGISTRATION NUMBER: 33,963  
REFERENCE/DOCKET NUMBER: 930049.407C5  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 206-622-4900

TELEFAX: 206-682-6031  
TELEX: 3723836  
INFORMATION FOR SEQ ID NO: 55:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 67 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
SEQUENCE DESCRIPTION: SEQ ID NO: 55:  
US-09-912-679-55

Query Match 40.6%; Score 52.4; DB 10; Length 67;  
Best Local Similarity 98.1%; Pred. No. 6.2e-10;  
Matches 53; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 72 TCGTGACGCAACCCCACTGCTGGGGCTTGSCCATAGGCAATCAGCGCATGCG 125  
|||  
DB 14 TCGTGACGCAACCCCACTGCTGGGGCTTGSCCATAGGCAATCAGCGCATGCG 67

Search completed: May 21, 2003, 04:39:45  
Job time: 122 secs





FT /note= "Inverted terminal repeat"  
 XX WO200127303-A1.  
 PN 19-APR-2001.  
 XX  
 PD 12-OCT-2000; 2000WO-US28221.  
 XX  
 PF 12-OCT-1999; 99US-0158780.  
 XX  
 PR (UYNC-) UNIV NORTH CAROLINA.  
 PA  
 XX Walsh CE, Chao H, Burstein H, Lynch CM, Stepan AM, Munson K;  
 PI WPI; 2001-273781/28.  
 DR P-PSDB; AAB67960.  
 DR  
 XX  
 PT New recombinant adeno-associated virus vector, useful for treating  
 PT haemophilia A, comprises heterologous nucleotide sequence encoding  
 PT B-domain deleted human factor VIII operably linked with liver-preferred  
 PT expression control element -  
 XX  
 XX PS Disclosure: Fig 6; 87pp; English.  
 XX  
 CC The specification describes a recombinant adeno-associated virus (rAAV)  
 CC vector. The vector comprises a heterologous nucleotide sequence  
 CC encoding B-domain deleted factor VIII operably linked with at least one  
 CC enhancer and at least one promoter. The method results in the production  
 CC of high titer rAAV vector stocks carrying the B-domain deleted factor  
 CC VIII transgenes and expression cassettes, which generate adequate titers  
 CC of virus for in vivo administration. The recombinant vectors are useful  
 CC for treating haemophilia A, where the liver expresses the encoded  
 CC B-domain deleted factor VIII, which is secreted into the blood. They are  
 CC also useful for the treatment of other coagulation disorders. The  
 CC present sequence encodes a B-domain deleted factor VIII.  
 XX  
 SQ Sequence 7914 BP; 2055 A; 1994 C; 1950 G; 1915 T; 0 other;  
 Query Match 100.0%; Score 129; DB 22; Length 7914;  
 Best Local Similarity 100.0%; Pred. No. 9.2e-35;  
 Matches 129; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTCCTTCTAAGTAACAGTACATGAACTTTACCCCGTGTGCTGGCAAGCGCTGTCTG 60  
 DB 150 CTCCTTCTAAGTAACAGTACATGAACTTTACCCCGTGTGCTGGCAAGCGCTGTCTG 209  
 QY 61 TGCCAAGTGTGCTGACCAACCCCACTGCTGGCGCTGGCAATAGGCATAGCGC 120  
 DB 210 TGCCAAGTGTGCTGACCAACCCCACTGCTGGCGCTGGCAATAGGCATAGCGC 269  
 QY 121 ATGCGGATC 129  
 DB 270 ATGCGGATC 278  
 RESULT 2  
 AAF84647  
 ID AAF84647 standard; DNA; 7944 BP.  
 XX  
 AC AAF84647;  
 XX  
 XX 29-JUN-2001 (first entry)  
 DT  
 XX Plasmid DL26 encoding human B-domain deleted factor VIII.  
 DE  
 XX Adeno-associated virus vector; B-domain; factor VIII; haemophilia A;  
 KW coagulation disorder; ss.  
 XX  
 OS Synthetic.  
 OS Homo sapiens.  
 OS Hepatitis B virus.  
 XX  
 FH Key Location/Qualifiers

FT misc-feature 1..146  
 FT /tag= a  
 FT /note= "Inverted terminal repeat"  
 FT enhancer 150..278  
 FT /tag= b  
 FT /note= "hepatitis B virus EnhI enhancer"  
 FT CDS 420..483  
 FT /tag= c  
 FT /note= "human B-domain deleted factor VIII"  
 FT polyA\_signal 4840..4914  
 FT /tag= d  
 FT /note= "TK polyA sequence"  
 FT misc-feature 4916..5084  
 FT /tag= e  
 FT /note= "Inverted terminal repeat"  
 XX  
 PN WO200127303-A1.  
 XX  
 PD 19-APR-2001.  
 XX  
 PF 12-OCT-2000; 2000WO-US28221.  
 XX  
 PR 12-OCT-1999; 99US-0158780.  
 XX  
 PA (UYNC-) UNIV NORTH CAROLINA.  
 XX  
 PI Walsh CE, Chao H, Burstein H, Lynch CM, Stepan AM, Munson K;  
 DR WPI; 2001-273781/28.  
 DR P-PSDB; AAB67959.  
 XX  
 XX New recombinant adeno-associated virus vector, useful for treating  
 PT haemophilia A, comprises heterologous nucleotide sequence encoding  
 PT B-domain deleted human factor VIII operably linked with liver-preferred  
 PT expression control element -  
 XX  
 XX PS Claim 64; Fig 1; 87pp; English.  
 XX  
 CC The specification describes a recombinant adeno-associated virus (rAAV)  
 CC vector. The vector comprises a heterologous nucleotide sequence  
 CC encoding B-domain deleted factor VIII operably linked with at least one  
 CC enhancer and at least one promoter. The method results in the production  
 CC of high titer rAAV vector stocks carrying the B-domain deleted factor  
 CC VIII transgenes and expression cassettes, which generate adequate titers  
 CC of virus for in vivo administration. The recombinant vectors are useful  
 CC for treating haemophilia A, where the liver expresses the encoded  
 CC B-domain deleted factor VIII, which is secreted into the blood. They are  
 CC also useful for the treatment of other coagulation disorders. The  
 CC present sequence encodes a B-domain deleted factor VIII.  
 XX  
 SQ Sequence 7944 BP; 2142 A; 1902 C; 1909 G; 1991 T; 0 other;  
 Query Match 100.0%; Score 129; DB 22; Length 7944;  
 Best Local Similarity 100.0%; Pred. No. 9.2e-35;  
 Matches 129; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 CTCCTTCTAAGTAACAGTACATGAACTTTACCCCGTGTGCTGGCAAGCGCTGTCTG 60  
 DB 150 CTCCTTCTAAGTAACAGTACATGAACTTTACCCCGTGTGCTGGCAAGCGCTGTCTG 209  
 QY 61 TGCCAAGTGTGCTGACCAACCCCACTGCTGGCGCTGGCAATAGGCATAGCGC 120  
 DB 210 TGCCAAGTGTGCTGACCAACCCCACTGCTGGCGCTGGCAATAGGCATAGCGC 269  
 QY 121 ATGCGGATC 129  
 DB 270 ATGCGGATC 278  
 RESULT 3  
 AAT73163  
 ID AAT73163 standard; cDNA; 587 BP.  
 XX

AC AAT73163;  
XX  
XX 08-APR-1998 (first entry)  
XX  
DE Post-translational regulatory element (PRE) of the Hepatitis B virus.  
XX  
XX Post-translational regulatory element; PRE; enhancer II; intronless gene;  
KW surface antigen gene; cytoplasmic accumulation; targeted delivery;  
KW near consensus splice sequence; blood coagulation factor; factor VIII;  
KW factor IX; ss.  
XX  
OS Hepatitis B virus.  
XX  
XX WO9733994-A1.  
XX  
XX 18-SEP-1997.  
XX  
XX 10-MAR-1997; 97WO-0503561.  
XX  
XX 11-MAR-1996; 96US-0683839.  
XX  
XX (IMMU-) IMMUNE RESPONSE CORP.  
XX  
XX Bidlingmeyer S, III CR;  
XX  
XX WPI; 1997-470874/43.  
XX  
XX Vector for increased expression of intronless genes - comprises  
PT intronless gene with at least one near consensus splice sequence, a  
PT promoter and at least one viral cis-acting post-transcriptional  
PT regulatory element  
XX  
XX  
PS Claim 3; Page 21; 59pp; English.  
XX  
XX The present sequence represents a post-translational regulatory element  
CC (PRE) of the Hepatitis B virus. This sequence encompasses enhancer II,  
CC and is within the transcribed portion of the surface antigen gene. This  
CC PRE sequence has been shown to function in cis to increase the  
CC steady-state levels of surface gene transcripts by facilitating  
CC cytoplasmic accumulation of these transcripts. The present PRE sequence  
CC was used to create a novel vector, comprising an intronless gene  
CC containing one or more near consensus splice sequences operably linked to a  
CC promoter sequence so that the gene is transcribed in a cell. One or more  
CC copies of a viral cis-acting PRE are also cloned into the vector, and are  
CC transcribed along with the gene, causing export of the gene transcript  
CC from the nucleus into the cytoplasm of the cell. The vector can be used  
CC to increase the expression of an intronless gene containing at least one  
CC near consensus splice sites, preferably cDNA encoding a blood coagulation  
CC factor, particularly Factor VIII or IX. The complex allows the targeted  
CC delivery of the vector to a specific cell, e.g. hepatocytes when the  
CC ligand is an asialoglycoprotein which binds the asialoglycoprotein  
CC receptor present on their surface.  
XX  
XX  
SQ Sequence 587 BP; 97 A; 199 C; 145 G; 146 T; 0 other;  
Query Match 95.3%; Score 123; DB 18; Length 587;  
Best Local Similarity 100.0%; Pred. No. 4,7e-33;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 3 CTTTCTAAGTAACAGTACATGACCTTTACCCGTTGCTGCGCAACGGCCTGCTGTG 62  
DB 2 CTTTCTAAGTAACAGTACATGACCTTTACCCGTTGCTGCGCAACGGCCTGCTGTG 61  
OY 63 CCAAGTGTGTTGCTGACGCAACCCCACTGGCTGGGGCTTGGCCATAGGCATCAGCGCAT 122  
DB 62 CCAAGTGTGTTGCTGACGCAACCCCACTGGCTGGGGCTTGGCCATAGGCATCAGCGCAT 121  
OY 123 GCG 125  
DB 122 GCG 124

RESULT 4

AAV69745  
ID AAV69745 standard; CDNA; 587 BP.  
XX  
XX AAV69745;  
XX  
XX 04-FEB-1999 (first entry)  
XX  
XX HBV post-transcriptional regulatory element (PRE) sequence.  
DE  
XX  
XX Hepatitis B virus; post-transcriptional regulatory element; PRE; HBV;  
KW viral transcript; ss.  
XX  
XX Hepatitis B virus.  
XX  
XX US5843770-A.  
XX  
XX 01-DEC-1998.  
XX  
XX 11-MAR-1996; 96US-0613861.  
XX  
XX 11-MAR-1996; 96US-0613861.  
XX  
XX (IMMU-) IMMUNE RESPONSE CORP.  
XX  
XX Gonzales JEN, III CR;  
XX  
XX WPI; 1999-044589/04.  
XX  
XX Hepatitis B virus antisense vector - directed against cis-acting  
PT post-transcriptional regulatory element  
XX  
XX  
PS Claim 1; Columns 13-14; 12pp; English.  
XX  
XX This sequence represents a hepatitis B virus cis-acting post-  
CC transcriptional regulatory element (PRE). The invention provides a vector  
CC encoding one or more antisense transcripts that are complementary to all  
CC or part of a HBV PRE where the PRE directs export of viral transcripts  
CC from the nucleus to the cytoplasm of a cell. A molecular complex  
CC comprising the above vector can be releasably linked to a conjugate of  
CC a nucleic acid binding agent and a ligand that binds to a component on  
CC the surface of a cell. The vector can be delivered to cells in vitro or  
CC in vivo to inhibit production of viruses having PRE sequences.  
XX  
XX  
SQ Sequence 587 BP; 97 A; 199 C; 145 G; 146 T; 0 other;  
Query Match 95.3%; Score 123; DB 20; Length 587;  
Best Local Similarity 100.0%; Pred. No. 4,7e-33;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 3 CTTTCTAAGTAACAGTACATGACCTTTACCCGTTGCTGCGCAACGGCCTGCTGTG 62  
DB 2 CTTTCTAAGTAACAGTACATGACCTTTACCCGTTGCTGCGCAACGGCCTGCTGTG 61  
OY 63 CCAAGTGTGTTGCTGACGCAACCCCACTGGCTGGGGCTTGGCCATAGGCATCAGCGCAT 122  
DB 62 CCAAGTGTGTTGCTGACGCAACCCCACTGGCTGGGGCTTGGCCATAGGCATCAGCGCAT 121  
OY 123 GCG 125  
DB 122 GCG 124

RESULT 5

AAH77169  
ID AAH77169 standard; DNA; 909 BP.  
XX  
XX AAH77169;  
XX  
XX 23-JAN-2002 (first entry)  
XX  
XX Regulatory and coding region of the X15 component in the X-myc construct.  
DE  
XX  
XX Transgenic mouse; cancer; oncogene; bicistronic hepatitis B virus; HBV;  
XX

KW X15-c-myc transgene; hepatocellular carcinoma; malignant liver tumour;  
XX X15; c-myc; murine; HBx; carcinogen; ds.  
OS Hepatitis B virus.  
XX US6274788-B1.  
XX 14-AUG-2001.  
XX 02-FEB-1999; 99US-0243282.  
XX 23-SEP-1998; 98IN-0002858.  
XX (TIGRE) INT CENT GENETIC ENG & BIOTECHNOLOGY.  
XX (NAIM) NAT INST IMMUNOLOGY.  
XX Kumar V, Singh M, Toley S, Anand R;  
XX WPI; 2002-009266/01.  
XX New bicistronic hepatitis B virus (HBV) X15-c-myc transgene, useful for  
XX producing transgenic mouse model systems for human hepatocellular  
XX carcinoma, comprises HBV X15 transgene and c-myc transgene -  
XX  
XX  
XX Claim 3; Fig 3; 12pp; English.  
XX  
XX This polynucleotide represents the sequence of the regulatory and coding  
XX regions of the X15 component in the X-myc construct. The invention  
XX relates to a bicistronic hepatitis B virus (HBV) X15-c-myc transgene,  
XX comprising of the HBV X15 gene and c-myc gene. The myc gene is known to  
XX be an activatable oncogene. The transgene encodes a truncated HBV X15  
XX protein that has amino acids 58-154 of HBV X15 and a murine c-myc  
XX construct is useful for screening a candidate substance (CS), to  
XX determine whether CS promotes hepatocellular carcinoma. This is  
XX determined by exposing a transgenic mouse to CS, and monitoring the mouse  
XX for the development of hepatocellular carcinoma, where an increase in the  
XX development of hepatocellular carcinoma in the transgenic mouse exposed  
XX to CS compared to the development of hepatocellular carcinoma in a  
XX transgenic mouse not exposed to CS, indicates that CS promotes  
XX hepatocellular carcinoma. The transgenic mice can be employed as a source  
XX for cell and tissue culture. The transgenic animal models comprising of  
XX the HBV X15-c-myc transgene for hepatocellular carcinoma are superior to  
XX any transgenic animal model system for hepatocellular carcinoma in that  
XX the transgenic mice develop more aggressive and accelerated onset of  
XX malignant liver tumours in all lobes causing death of the affected  
XX animals in 20-22 weeks, that is faster than the time taken by the other  
XX transgenic animals to even develop a tumour.  
XX  
XX Sequence 909 BP; 210 A; 236 C; 211 G; 252 T; 0 other;  
SQ  
Query Match 95.3%; Score 123; DB 24; Length 909;  
Best Local Similarity 100.0%; Pred. No. 5.5e-33;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 3 CTTTCTAAGTAACAGTACATGACCTTACCCCGTTGCTGGCAACGGCGCTGTGTG 62  
DB 286 CTTTCTAAGTAACAGTACATGACCTTACCCCGTTGCTGGCAACGGCGCTGTGTG 345  
OY 63 CCAAGTGTGGTGTGACGCAACCCCACTGGGCTGGGCAATGAGGCATGAGGCAT 122  
DB 346 CCAAGTGTGGTGTGACGCAACCCCACTGGGCTGGGCAATGAGGCATGAGGCAT 405  
OY 123 GCG 125  
DB 406 GCG 408  
RESULT 6  
AA223281  
ID AA223281 standard; DNA; 3221 BP.  
XX  
AC AA223281;

XX 31-JAN-2000 (first entry)  
DT DNA sequence of the genome of HBV adv 2.  
XX  
DE  
XX Hepatitis B virus; HBV; recombinant; pol gene; X gene; surface antigen;  
XX liver; anti-viral; anti-tumor; gene therapy; single-gene defect;  
XX genetic disorder; familial hypercholesterolemia; neoplastic gene;  
XX ornithine transcarbamylase deficiency; ss.  
XX  
XX Hepatitis b virus.  
XX  
XX US5981274-A.  
XX 09-NOV-1999.  
XX  
XX 18-SEP-1996; 96US-0715808.  
XX 18-SEP-1996; 96US-0715808.  
XX 18-SEP-1996; 96US-0715808.  
XX  
XX (CHAI/) CHAISOMCHIT S.  
XX (CHAI/) CHANG L.  
XX (TYRR/) TYRRELL D L J.  
XX  
XX Chang L, Chaisomchit S, Tyrrell DLJ;  
XX WPI; 1999-633330/54.  
XX  
XX Recombinant hepatitis B virus genome containing heterologous gene  
XX sequences useful for treating liver infections -  
XX  
XX Disclosure; Columns 35-39; 53pp; English.  
XX  
XX The invention relates to a recombinant hepatitis B virus genome (HBV)  
XX that comprises heterologous gene sequences which express at least one  
XX functional heterologous gene product. A host cell transfected with a  
XX recombinant HBV genome comprising pol gene sequences, X gene sequences  
XX and surface antigen gene (preS1/preS2/S gene) sequences and heterologous  
XX gene sequences can be used to express at least one functional  
XX heterologous gene product. The invention also provides a method for  
XX encapsulating a recombinant HBV genome. The recombinant HBV genomes are  
XX useful for the expression of functional heterologous gene products in  
XX liver cells. The vectors can be used for anti-viral, anti-tumor and/or  
XX gene therapy and particularly for the correction of inherited single-gene  
XX defects. Human genetic disorders which can be treated by expression of  
XX missing or mutant genes in the liver are familial hypercholesterolemia  
XX and ornithine transcarbamylase deficiency. Primary tumors of the liver  
XX may benefit from the expression of anti-neoplastic genes in the liver.  
XX Existing retroviral vectors and other animal viruses which are used to  
XX deliver foreign genes are not liver-specific with regard to their  
XX infection or expression unlike hepatitis B viral vectors. Human hepatitis  
XX B virus can be delivered through the circulation so there is no  
XX requirement for tissue culture for ex vivo liver-directed gene therapy.  
XX The present sequence represents the DNA sequence of the genome of HBV  
XX adv 2 which comprises the pol gene sequences, X gene sequences and  
XX surface antigen gene (preS1/preS2/S gene) sequences.  
XX  
XX Sequence 3221 BP; 740 A; 869 G; 708 G; 904 T; 0 other;  
SQ  
Query Match 95.3%; Score 123; DB 20; Length 3221;  
Best Local Similarity 100.0%; Pred. No. 8.4e-33;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
OY 3 CTTTCTAAGTAACAGTACATGACCTTACCCCGTTGCTGGCAACGGCGCTGTGTG 62  
DB 1118 CTTTCTAAGTAACAGTACATGACCTTACCCCGTTGCTGGCAACGGCGCTGTGTG 1177  
OY 63 CCAAGTGTGGTGTGACGCAACCCCACTGGGCTGGGCAATGAGGCATGAGGCAT 122  
DB 1178 CCAAGTGTGGTGTGACGCAACCCCACTGGGCTGGGCAATGAGGCATGAGGCAT 1237  
OY 123 GCG 125  
DB 111

ID 1238 GCG 1240

RESULT 7  
AADI4316  
ID AADI4316 standard; DNA; 4084 BP.  
XX  
AC AADI4316;  
XX  
XX 06-NOV-2001 (first entry)  
DE Hepatitis B virus (HBV) 1.28 genome.  
XX  
XX Hepatitis B virus; HBV; altered sensitivity; agent; detection;  
KM PCR primer; Hepatitis B surface antigen; HbsAg; ds.  
XX  
OS Hepatitis B virus.  
XX  
PN W0200157244-A1.  
XX  
PD 09-AUG-2001.  
XX  
XX 02-FEB-2001; 2001WO-AU00098.  
XX  
XX 03-FEB-2000; 2000US-0179948.  
XX  
XX (MELB-) MELBOURNE HEALTH.  
PA (PENN-) PENN STATE RES FOUND.  
XX  
PI Delaney W, Locarnini SA, Chen RYM, Bartholomeusz A, Isom H;  
DR WPI: 2001-496926/54.  
XX  
XX Detecting hepatitis B virus variant with altered sensitivity to agent,  
PT comprises infecting genetic construct containing replication competent  
PT genome to cells, contacting cells with agent and detecting replication  
PT of variant -  
XX  
XX Example 7; Fig 5A; 110pp; English.  
XX  
XX The invention relates to a method of detecting variant hepatitis B virus  
CC (HBV) which exhibits altered sensitivity to agents. The method involves  
CC infecting a genetic construct containing a replication competent amount  
CC of the genome from variant HBV contained in or fused to a baculovirus  
CC genome; contacting cells with the agent to be tested; culturing cells  
CC under conditions sufficient for the variant HBV to replicate; express  
CC genetic sequences, and/or assemble, and/or release viral particles; and  
CC determining replication of variant HBV using viral component-detection  
CC means. The method is useful for detecting variant HBV which exhibits  
CC altered sensitivity to agents. The present sequence is the HBV 1.28  
CC genome.  
XX  
XX Sequence 4084 BP; 920 A; 1105 C; 914 G; 1145 T; 0 other;  
SO

Query Match 95.3%; Score 123; DB 22; Length 4084;  
Best Local Similarity 100.0%; Pred. No. 9.1e-33;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 CTTTGAAGTAACAGTACATGACCTTACCCCGTGTGCTGCGCAACGGCCGTGCTGTG 62  
DB 2886 CTTTGAAGTAACAGTACATGACCTTACCCCGTGTGCTGCGCAACGGCCGTGCTGTG 2945

QY 63 CCAAGTGTGCTGACGCAACCCCACTGCGTGGGGCTTGGCCATAGGCCATCAGCGCAT 122  
DB 2946 CCAAGTGTGCTGACGCAACCCCACTGCGTGGGGCTTGGCCATAGGCCATCAGCGCAT 3005

QY 123 GCG 125  
DB 3006 GCG 3008

RESULT 8  
AADI4317

ID AADI4317 standard; DNA; 4496 BP.  
XX  
XX AADI4317;  
XX  
XX 06-NOV-2001 (first entry)  
DE Hepatitis B virus (HBV) 1.5 genome.  
XX  
XX Hepatitis B virus; HBV; altered sensitivity; agent; detection;  
KM PCR primer; Hepatitis B surface antigen; HbsAg; ds.  
XX  
XX Hepatitis B virus.  
XX  
PN W0200157244-A1.  
XX  
PD 09-AUG-2001.  
XX  
XX 02-FEB-2001; 2001WO-AU00098.  
XX  
XX 03-FEB-2000; 2000US-0179948.  
XX  
XX (MELB-) MELBOURNE HEALTH.  
PA (PENN-) PENN STATE RES FOUND.  
XX  
PI Delaney W, Locarnini SA, Chen RYM, Bartholomeusz A, Isom H;  
DR WPI: 2001-496926/54.  
XX  
XX Detecting hepatitis B virus variant with altered sensitivity to agent,  
PT comprises infecting genetic construct containing replication competent  
PT genome to cells, contacting cells with agent and detecting replication  
PT of variant -  
XX  
XX Example 7; Fig 5B; 110pp; English.  
XX  
XX The invention relates to a method of detecting variant hepatitis B virus  
CC (HBV) which exhibits altered sensitivity to agents. The method involves  
CC infecting a genetic construct containing a replication competent amount  
CC of the genome from variant HBV contained in or fused to a baculovirus  
CC genome; contacting cells with the agent to be tested; culturing cells  
CC under conditions sufficient for the variant HBV to replicate; express  
CC genetic sequences, and/or assemble, and/or release viral particles; and  
CC determining replication of variant HBV using viral component-detection  
CC means. The method is useful for detecting variant HBV which exhibits  
CC altered sensitivity to agents. The present sequence is the HBV 1.5  
CC genome.  
XX  
XX Sequence 4496 BP; 999 A; 1229 C; 1011 G; 1257 T; 0 other;  
SO

Query Match 95.3%; Score 123; DB 22; Length 4496;  
Best Local Similarity 100.0%; Pred. No. 9.5e-33;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 CTTTGAAGTAACAGTACATGACCTTACCCCGTGTGCTGCGCAACGGCCGTGCTGTG 62  
DB 77 CTTTGAAGTAACAGTACATGACCTTACCCCGTGTGCTGCGCAACGGCCGTGCTGTG 136

QY 63 CCAAGTGTGCTGACGCAACCCCACTGCGTGGGGCTTGGCCATAGGCCATCAGCGCAT 122  
DB 137 CCAAGTGTGCTGACGCAACCCCACTGCGTGGGGCTTGGCCATAGGCCATCAGCGCAT 196

QY 123 GCG 125  
DB 197 GCG 199

RESULT 9  
AAV69746/C  
ID AAV69746 standard; cDNA; 4525 BP.  
XX  
XX AAV69746;  
XX  
XX 04-FEB-1999 (first entry)

```

XX DE Nucleotide sequence of one copy antisense plasmid pCMVasPRE-RZ.
XX XX Hepatitis B virus; post-transcriptional regulatory element; PRE; HBV;
XX KW viral transcript; pCMVasPRE-RZ; ss.
XX OS Synthetic.
XX OS Hepatitis b virus.
XX XX US5843770-A.
XX XX 01-DEC-1998.
XX PD 11-MAR-1996; 96US-0613861.
XX PF 11-MAR-1996; 96US-0613861.
XX PR 11-MAR-1996; 96US-0613861.
XX PA (IMMU-) IMMUNE RESPONSE CORP.
XX PI Gonzales JEN, Ili CR;
XX DR WPI; 1999-044589/04.
XX PT Hepatitis B virus antisense vector - directed against cis-acting
XX PT post-transcriptional regulatory element
XX PS Claim 3; Columns 13-18; 12pp; English.
XX CC This represents the nucleotide sequence of an one copy antisense plasmid
XX CC pCMVasPRE-RZ. The plasmid contains a hepatitis B virus cis-acting post-
XX CC transcriptional regulatory element (PRE). The invention provides such a
XX CC vector encoding one or more antisense transcripts that are complementary
XX CC to all or part of a HBV PRE where the PRE directs export of viral
XX CC transcripts from the nucleus to the cytoplasm of a cell. A molecular
XX CC complex comprising the above vector can be releasably linked to a
XX CC conjugate of a nucleic acid binding agent and a ligand that binds to a
XX CC component on the surface of a cell. The vector can be delivered to cells
XX CC in vitro or in vivo to inhibit production of viruses having PRE
XX CC sequences.
SQ Sequence 4525 BP; 1115 A; 1132 C; 1168 G; 1110 T; 0 other;

Query Match 95.3%; Score 123; DB 20; Length 4525;
Best Local Similarity 100.0%; Pred. No. 9.5e-33;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 CTTTCTAAGTAAACAGTACATGAACTTTACCCCGTTGCTGGCAACGGCGCTGCTGTG 62
DB 3727 CTTTCTAAGTAAACAGTACATGAACTTTACCCCGTTGCTGGCAACGGCGCTGCTGTG 3668

QY 63 CCAAGTGTGTTGCTGACGCAACCCCACTGCTGGGCTTGGCCATAGGCCATCAGCGCAT 122
DB 3667 CCAAGTGTGTTGCTGACGCAACCCCACTGCTGGGCTTGGCCATAGGCCATCAGCGCAT 3608

QY 123 GCG 125
DB 3607 GCG 3605

RESULT 10
AA23285
ID AA23285 standard; DNA; 6371 BP.
XX AC AA23285;
XX XX 31-JAN-2000 (first entry)
XX DE DNA sequence of plasmid pTHBVT.
XX XX Hepatitis B virus; HBV; recombinant; pol gene; X gene; surface antigen;
XX KW liver; anti-viral; anti-tumor; gene therapy; single-gene defect;
XX KW genetic disorder; familial hypercholesterolemia; neoplastic gene;
XX KW ornithine transcarbamylase deficiency; ss.

```

```

XX OS Synthetic.
XX OS Hepatitis b virus.
XX XX US5981274-A.
XX XX 09-NOV-1999.
XX PF 18-SEP-1996; 96US-0715808.
XX PR 18-SEP-1996; 96US-0715808.
XX PA (CHAI/) CHAISOMCHIT S.
XX PA (CHAN/) CHANG L.
XX PA (TYRR/) TYRRELL D L J.
XX PI Chang L, Chaisomchit S, Tyrrell DLJ;
XX DR WPI; 1999-633330/54.
XX PT Recombinant hepatitis B virus genome containing heterologous gene
XX PT sequences useful for treating liver infections -
XX PS Example 1; Columns 47-54; 53pp; English.
XX CC The invention relates to a recombinant hepatitis B virus genome (HBV)
XX CC that comprises heterologous gene sequences which express at least one
XX CC functional heterologous gene product. A host cell transfected with a
XX CC recombinant HBV genome comprising pol gene sequences, X gene sequences
XX CC and surface antigen gene (pres1/pres2/5 gene) sequences and heterologous
XX CC gene sequences can be used to express at least one functional
XX CC heterologous gene product. The invention also provides a method for
XX CC encapsulating a recombinant HBV genome. The recombinant HBV genomes are
XX CC useful for the expression of functional heterologous gene products in
XX CC liver cells. The vectors can be used for anti-viral, anti-tumor and/or
XX CC gene therapy and particularly for the correction of inherited single-gene
XX CC defects. Human genetic disorders which can be treated by expression of
XX CC missing or mutant genes in the liver are familial hypercholesterolemia
XX CC and ornithine transcarbamylase deficiency. Primary tumors of the liver
XX CC may benefit from the expression of anti-neoplastic genes in the liver.
XX CC Existing retroviral vectors and other animal viruses which are used to
XX CC deliver foreign genes are not liver-specific with regard to their
XX CC infection or expression unlike hepatitis B viral vectors. Human hepatitis
XX CC B virus can be delivered through the circulation so there is no
XX CC requirement for tissue culture for ex vivo liver-directed gene therapy.
XX CC The present sequence represents the DNA sequence of the plasmid pTHBVT
XX CC which comprises HBV sequences.
SQ Sequence 6371 BP; 1568 A; 1650 C; 1485 G; 1668 T; 0 other;

Query Match 95.3%; Score 123; DB 20; Length 6371;
Best Local Similarity 100.0%; Pred. No. 1.1e-32;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 3 CTTTCTAAGTAAACAGTACATGAACTTTACCCCGTTGCTGGCAACGGCGCTGCTGTG 62
DB 1364 CTTTCTAAGTAAACAGTACATGAACTTTACCCCGTTGCTGGCAACGGCGCTGCTGTG 1423

QY 63 CCAAGTGTGTTGCTGACGCAACCCCACTGCTGGGCTTGGCCATAGGCCATCAGCGCAT 122
DB 1424 CCAAGTGTGTTGCTGACGCAACCCCACTGCTGGGCTTGGCCATAGGCCATCAGCGCAT 1483

QY 123 GCG 125
DB 1484 GCG 1486

RESULT 11
AA23292
ID AA23292 standard; DNA; 6371 BP.
XX AC AA23292;
XX XX

```

31-JAN-2000 (first entry)

DNA sequence of plasmid pTHBVTX-.

Hepatitis B virus; HBV; recombinant; pol gene; X gene; surface antigen; liver; anti-viral; anti-tumor; gene therapy; single-gene defect; genetic disorder; familial hypercholesterolemia; neoplastic gene; ornithine transcarbamylase deficiency; ss.

Synthetic.

Hepatitis b virus.

US5981274-A.

09-NOV-1999.

18-SEP-1996; 96US-0715808.

18-SEP-1996; 96US-0715808.

18-SEP-1996; 96US-0715808.

(CHAI/) CHAISOMCHIT S.

(CHAN/) CHANG L.

(TYRR/) TYRRELL D L J.

Chang L, Chaisomchit S, Tyrrell DLJ;

WPI: 1999-633330/54.

Recombinant hepatitis B virus genome containing heterologous gene sequences useful for treating liver infections -

Example 2; Columns 63-70; 53pp; English.

The invention relates to a recombinant hepatitis B virus genome (HBV) that comprises heterologous gene sequences which express at least one functional heterologous gene product. A host cell transfected with a recombinant HBV genome comprising pol gene sequences, X gene sequences and surface antigen gene (pres1/pres2/S gene) sequences and heterologous gene sequences can be used to express at least one functional heterologous gene product. The invention also provides a method for encapsidating a recombinant HBV genome. The recombinant HBV genomes are useful for the expression of functional heterologous gene products in liver cells. The vectors can be used for anti-viral, anti-tumor and/or gene therapy and particularly for the correction of inherited single-gene defects. Human genetic disorders which can be treated by expression of missing or mutant genes in the liver are familial hypercholesterolemia and ornithine transcarbamylase deficiency. Primary tumors of the liver may benefit from the expression of anti-neoplastic genes in the liver. Existing retroviral vectors and other animal viruses which are used to deliver foreign genes are not liver-specific with regard to their infection or expression unlike hepatitis B viral vectors. Human hepatitis B virus can be delivered through the circulation so there is no requirement for tissue culture for ex vivo liver-directed gene therapy. The present sequence represents the DNA sequence of the plasmid pTHBVTX-.

Sequence 6371 BP; 1567 A; 1649 C; 1485 G; 1670 T; 0 other;

Query Match 95.3%; Score 123; DB 20; Length 6371;

Best Local Similarity 100.0%; Pred. No. 1.1e-32;

Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

3 CTTTCTAGTAAGAGTACAGTACGCTTACCGCTGCTGCGCAACGCGCTGCTGTG 62

1364 CTTTCTAGTAAGAGTACAGTACGCTTACCGCTGCTGCGCAACGCGCTGCTGTG 1423

63 CCAAGTGTGCTGACGCAACCCCACTGCGCTTGGCCATAGGCGCATACCGCAT 122

1424 CCAAGTGTGCTGACGCAACCCCACTGCGCTTGGCCATAGGCGCATACCGCAT 1483

123 GCG 125

1484 GCG 1486

RESULT 12

AA23294

ID AA23294 standard; DNA; 6375 BP.

XX AA23294;

AC AA23294;

XX

31-JAN-2000 (first entry)

DNA sequence of plasmid pTHBVTX-.

Hepatitis B virus; HBV; recombinant; pol gene; X gene; surface antigen; liver; anti-viral; anti-tumor; gene therapy; single-gene defect; genetic disorder; familial hypercholesterolemia; neoplastic gene; ornithine transcarbamylase deficiency; ss.

Synthetic.

Hepatitis b virus.

US5981274-A.

09-NOV-1999.

18-SEP-1996; 96US-0715808.

18-SEP-1996; 96US-0715808.

18-SEP-1996; 96US-0715808.

(CHAI/) CHAISOMCHIT S.

(CHAN/) CHANG L.

(TYRR/) TYRRELL D L J.

Chang L, Chaisomchit S, Tyrrell DLJ;

WPI: 1999-633330/54.

Recombinant hepatitis B virus genome containing heterologous gene sequences useful for treating liver infections -

Example 3; Columns 75-82; 53pp; English.

The invention relates to a recombinant hepatitis B virus genome (HBV) that comprises heterologous gene sequences which express at least one functional heterologous gene product. A host cell transfected with a recombinant HBV genome comprising pol gene sequences, X gene sequences and surface antigen gene (pres1/pres2/S gene) sequences and heterologous gene sequences can be used to express at least one functional heterologous gene product. The invention also provides a method for encapsidating a recombinant HBV genome. The recombinant HBV genomes are useful for the expression of functional heterologous gene products in liver cells. The vectors can be used for anti-viral, anti-tumor and/or gene therapy and particularly for the correction of inherited single-gene defects. Human genetic disorders which can be treated by expression of missing or mutant genes in the liver are familial hypercholesterolemia and ornithine transcarbamylase deficiency. Primary tumors of the liver may benefit from the expression of anti-neoplastic genes in the liver. Existing retroviral vectors and other animal viruses which are used to deliver foreign genes are not liver-specific with regard to their infection or expression unlike hepatitis B viral vectors. Human hepatitis B virus can be delivered through the circulation so there is no requirement for tissue culture for ex vivo liver-directed gene therapy. The present sequence represents the DNA sequence of the plasmid pTHBVTX-.

Sequence 6375 BP; 1568 A; 1652 C; 1487 G; 1668 T; 0 other;

Query Match 95.3%; Score 123; DB 20; Length 6375;

Best Local Similarity 100.0%; Pred. No. 1.1e-32;

Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

3 CTTTCTAGTAAGAGTACAGTACGCTTACCGCTGCTGCGCAACGCGCTGCTGTG 62

1364 CTTTCTAGTAAGAGTACAGTACGCTTACCGCTGCTGCGCAACGCGCTGCTGTG 1423

63 CCAAGTGTGCTGACGCAACCCCACTGCGCTTGGCCATAGGCGCATACCGCAT 122

Db	1424	CAGAGTGTTCGACGAAACCCCACTGGCTGGGCGTTGCCATATGCGCAT	1483
Oy	123	GCG 125 	
Dd	1484	GCG 1486	
RESULT 13			
ID	AA223282	standard; DNA; 9325 BP.	
XX	AA223282;		
XX	31-JAN-2000	(first entry)	
DE	DNA sequence of plasmid pTHBV-d.		
XX	Hepatitis B virus; HBV; recombinant; pol gene; X gene; surface antigen;		
RW	liver; anti-viral; anti-tumor; gene therapy; single-gene defect;		
KW	genetic disorder; familial hypercholesterolemia; neoplastic gene;		
KM	ornithine transcarbamylase deficiency; ss.		
XX	Synthetic.		
OS	Hepatitis D virus.		
XX	US5981274-A.		
PN	09-NOV-1999.		
PD	18-SEP-1996;	96US-0715808.	
PF	18-SEP-1996;	96US-0715808.	
XX	18-SEP-1996;	96US-0715808.	
PR	(CHAI/) CHAISOMCHIT S.		
PA	(CHAN/) CHANG L.		
PA	(TYRR/) TYRRELL D L J.		
XX	Chang L, Chaisomchit S, Tyrrell DLJ;		
PI	WPt; 1999-633330/54.		
DR	Recombinant hepatitis B virus genome containing heterologous gene sequences useful for treating liver infections -		
PT	Example 1; Columns 39-48; 53pp; English.		
PS	The invention relates to a recombinant hepatitis B virus genome (HBV) that comprises heterologous gene sequences which express at least one functional heterologous gene product. A host cell transfected with a recombinant HBV genome comprising pol gene sequences, X gene sequences and surface antigen gene (pres1/pres2/5 gene) sequences and heterologous gene sequences can be used to express at least one functional heterologous gene product. The invention also provides a method for encapsulating a recombinant HBV genome. The recombinant HBV genomes are useful for the expression of functional heterologous gene products in liver cells. The vectors can be used for anti-viral, anti-tumor and/or gene therapy and particularly for the correction of inherited single-gene defects. Human genetic disorders which can be treated by expression of missing or mutant genes in the liver are familial hypercholesterolemia and ornithine transcarbamylase deficiency. Primary tumors of the liver may benefit from the expression of anti-neoplastic genes in the liver. Existing retroviral vectors and other animal viruses which are used to deliver foreign genes are not liver-specific with regard to their infection or expression unlike hepatitis B viral vectors. Human hepatitis B virus can be delivered through the circulation so there is no requirement for tissue culture for ex vivo liver-directed gene therapy. The present sequence represents the DNA sequence of the plasmid pTHBV-d which comprises HBV sequences.		
CC	Sequence 9325 BP; 2227 A; 2448 C; 2132 G; 2518 T; 0 other;		
CC	Query Match	95.3%; Score 123; DB 20; Length 9325;	

QY	3	CTTCTTAAGTAAACAGTACATGAAACCTTTACCCGTTGCTCGGCAACGGCTGTCTGTG	62
Db	1364	CTTCTTAAGTAAACAGTACATGAAACCTTTACCCGTTGCTCGGCAACGGCTGTCTGTG	1423
OY	63	CCAAAGTCTTCTGACGCAACCCCACTGCTGCGGCTTGCCATFAGGCCATCAGCGCAT	122
Db	1424	CCAAAGTCTTCTGACGCAACCCCACTGCTGCGGCTTGCCATFAGGCCATCAGCGCAT	1483
OY	123	GCG 125	
Db	1484	GCG 1486	
RESULT 14			
AA73164			
ID	AA73164	standard; cDNA; 9354 BP.	
XX	AA73164;		
XX	08-APR-1998	(first entry)	
XX	CDNA encoding human B-domain deleted factor VIII.		
XX	Post-translational regulatory element; PRE; enhancer II; intronless gene;		
XX	surface antigen gene; cytoplasmic accumulation; targeted delivery;		
KW	near consensus splice sequence; blood coagulation factor; factor VIII;		
KX	factor IX, ss.		
XX	Homo sapiens.		
OS			
XX	Key	Location/Qualifiers	
XX	CDS	2965..7380	
FT	misc_feature	5165..5174	
FT	misc_feature	5695..5703	
FT	misc_feature	6320..6328	
FT	misc_feature	7045..7053	
FT	misc_feature	7143..7152	
FT	misc_feature	3296..3312	
FT	misc_feature	4798..4817	
FT	misc_feature	5333..5355	
FT	misc_feature	5520..5538	
FT	misc_feature	5604..5632	
FT	misc_feature	5717..5745	



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FT FT /*tag= n /note= "3', near consensus site"
FT FT misc-feature 6239..6258
FT FT /*tag= o /note= "3', near consensus site"
FT FT misc-feature 6658..6682
FT FT /*tag= p /note= "3', near consensus site"
FT FT misc-feature 7159..7176
FT FT /*tag= q /note= "3', near consensus site"
FT FT misc-feature 7196..7209
FT FT /*tag= r /note= "3', near consensus site"
FT FT misc-feature 7289..7315
FT FT /*tag= s /note= "3', near consensus site"
FT FT misc-feature 7411..7429
FT FT /*tag= t /note= "3', near consensus site"
FT FT misc-feature 7611..8197
FT FT /*tag= u /note= "PRE sequence"
XX XX WO9733994-A1.
XX PD 18-SEP-1997.
XX PF 10-MAR-1997; 97WO-US03561.
XX PR 11-MAR-1996; 96US-0683839.
XX PA (IMMU-) IMMUNE RESPONSE CORP.
XX BIddingmaier S, ILL CR;
XX DR WPI: 1997-470874/43.
XX DR P-PSDB; AAMW23414.
XX PT Vector for increased expression of intronless genes - comprises
XX PT intronless gene with at least one near consensus splice sequence, a
XX PT promoter and at least one viral cis-acting post-transcriptional
XX PT regulatory element
XX EX Example 1; Pages 21-31; 59pp; English.
XX PS The present sequence represents human B-domain deleted factor VIII
XX CC cDNA, and a post-translational regulatory element (PRE) of the
XX CC Hepatitis B virus, which is present 3' of the STOP codon for Factor VIII.
XX CC PRE sequences have been shown to function in cis to increase the
XX CC steady-state levels of surface gene transcripts by facilitating
XX CC cytoplasmic accumulation of these transcripts. The present sequence
XX CC is part of a novel vector, comprising an intronless gene containing
XX CC 1 or more near consensus splice sequences operably linked to a
XX CC promoter sequence so that the gene is transcribed in a cell.
XX CC Intronless gene transcripts which contain near consensus splice site
XX CC sequences are believed to get tied up in the nucleus of the cell where
XX CC splicing occurs, rather than being transported to the cytoplasm where
XX CC they can be translated into proteins. The PRE sequences are transcribed
XX CC along with the gene, causing export of the gene transcript from the
XX CC nucleus into the cytoplasm of the cell. The vector can be used
XX CC to increase the expression of an intronless gene containing at least one
XX CC near consensus splice sites, preferably cDNA encoding a blood coagulation
XX CC factor, particularly Factor VIII or IX. The complex allows the targeted
XX CC delivery of the vector to a specific cell, e.g. hepatocytes when the
XX CC ligand is an asialoglycoprotein which binds the asialoglycoprotein
XX CC receptor present on their surface.
XX SQ Sequence 9354 BP; 2506 A; 2239 C; 2161 G; 2448 T; 0 other;
Query Match 95.3%; Score 123; DB 18; Length 9354;
Best Local Similarity 100.0%; Pred. No. 1,2e-32;
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0

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QY	3	CTTTCAAGATTAACACATGATCAATGAACCTTTACCCCGTTGCGGCAACGGCCTGGTCTGTG	62
Db	7612	CTTTCAAGATTAACACATGATCAATGAACCTTTACCCCGTTGCGGCAACGGCCTGGTCTGTG	7611
QY	63	CCAAGTGTGTTGCTGACGCCAACCCCACTGGCTGGGGCTTGGCCATAGCCATCAGCCGAT	122
Db	7672	CCAAGTGTGTTGCTGACGCCAACCCCACTGGCTGGGGCTTGGCCATAGCCATCAGCCGAT	7731
QY	123	GCG 125	
Db	7732	GCG 7734	
RESULT 15			
ID	AAZ23286	AAZ23286 standard; DNA; 9859 BP.	
XX	AAZ23286;		
XX	31-JAN-2000	(first entry)	
DE	DNA sequence of plasmid pTHEVT-d.		
XX	Hepatitis B virus; HBV; recombinant; pol gene; X gene; surface antigen;		
KW	liver; anti-viral; anti-tumor; gene therapy; single-gene defect;		
KW	genetic disorder; familial hypercholesterolemia; neoplastic gene;		
KW	ornithine transcarbamylase deficiency; ss.		
XX	Synthetic.		
OS	Hepatitis b virus.		
XX	US5981274-A.		
PN	09-NOV-1999.		
PD	18-SEP-1996; 96US-0715808.		
PF	18-SEP-1996; 96US-0715808.		
PR	18-SEP-1996; 96US-0715808.		
XX	(CHARI/) CHARISOMCHT S.		
PA	(CHAN/) CHANG L.		
PA	(TYRR/) TYRRELL D L J.		
XX	Chang L, Chaisomchit S, Tyrrell DLJ;		
PI	WPI: 1999-633330/54.		
DR	Recombinant hepatitis B virus genome containing heterologous gene		
PT	sequences useful for treating liver infections -		
PT	Example 1; Columns 53-62; 53pp; English.		
XX	The invention relates to a recombinant hepatitis B virus genome (HBV)		
CC	that comprises heterologous gene sequences which express at least one		
CC	functional heterologous gene product. A host cell transfected with a		
CC	recombinant HBV genome comprising pol gene sequences, X gene sequences		
CC	and surface antigen gene (pres1/pres2/S gene) sequences and heterologous		
CC	gene sequences can be used to express at least one functional		
CC	heterologous gene product. The invention also provides a method for		
CC	enhancing/expressing a recombinant HBV genome. The recombinant HBV genomes are		
CC	useful for the expression of functional heterologous gene products in		
CC	liver cells. The vectors can be used for anti-viral, anti-tumor and/or		
CC	gene therapy and particularly for the correction of inherited single-gene		
CC	defects. Human genetic disorders which can be treated by expression of		
CC	missing or mutant genes in the liver are familial hypercholesterolemia		
CC	and ornithine transcarbamylase deficiency. Primary tumors of the liver		
CC	may benefit from the expression of anti-neoplastic genes in the liver.		
CC	Existing retroviral vectors and other animal viruses which are used to		
CC	deliver foreign genes are not liver-specific with regard to their		
CC	infection or expression unlike hepatitis B viral vectors. Human hepatitis		
CC	B virus can be delivered through the circulation so there is no		
CC	requirement for tissue culture for ex vivo liver-directed gene therapy.		

CC The present sequence represents the DNA sequence of the plasmid pTHRTV-d.  
XX  
SQ Sequence 9859 BP; 2389 A; 2590 C; 2254 G; 2626 T; 0 other;

Query Match 95.3%; Score 123; DB 20; Length 9859;  
Best Local Similarity 100.0%; Pred. No. 1.2e-32;  
Matches 123; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

OY 3 CTTCTAAGTAAACAGTACATGACCTTTACCCGTTGCTCGGCACGCGCTGCTGTG 62  
|||  
DB 1364 CTTCTAAGTAAACAGTACATGACCTTTACCCGTTGCTCGGCACGCGCTGCTGTG 1423  
OY 63 CCAAGTGTGCTGACGCAACCCCACTGGCTGGGCTTGCCATAGCCATCAGCGCAT 122  
|||  
DB 1424 CCAAGTGTGCTGACGCAACCCCACTGGCTGGGCTTGCCATAGCCATCAGCGCAT 1483  
OY 123 GCG 125  
|||  
DB 1484 GCG 1486

Search completed: May 21, 2003, 03:56:18  
Job time : 223 secs